



STIC Search Report

Biotech-Chem Library

STIC Database Tracking Number: 168136

TO: Jeanine Goldberg
Location: rem/2D15/2C70
Art Unit: 1634
Wednesday, July 13, 2005
Case Serial Number: 10/681199

From: Toby Port
Location: Biotech-Chem Library
REM-1A59
Phone: 571-272-2523

toby.port@uspto.gov

Search Notes

Examiner Goldberg,

See attached results.

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Thank you for using STIC search services!

Toby Port
X22523

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STIC-Biotech/ChemLib

158136

mg

From: Goldberg, Jeanine
Sent: Tuesday, July 05, 2005 6:20 AM
To: STIC-Biotech/ChemLib
Subject: 10/681199- Search

Please search SEQ ID NO: 1 in all databases.
Please also do a mer search of SEQ ID NO: 1.

THANK YOU

Jeanine Anne Goldberg
1634
571-272-0743
REM 2D15

2070

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Searcher: _____
Searcher Phone: 2-_____
Date Searcher Picked up: _____
Date Completed: _____
Searcher Prep/Rev. Time: _____
Online Time: _____

Type of Search

NA#: _____ AA#: _____
Interference: _____ SPDI: _____
S/L: _____ Oligomer: _____
Encode/Transl: _____
Structure#: _____ Text: _____
Inventor: _____ Litigation: _____

Vendors and cost where applicable

STN: _____
DIALOG: _____
QUESTEL/ORBIT: _____
LEXIS/NEXIS: _____
SEQUENCE SYSTEM: _____
WWW/Internet: _____
Other(Specify): _____

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OM nucleic - nucleic search, using sw model
Run on: July 8, 2005, 04:46:25 ; Search time 758 Seconds
(without alignments)
9863.632 Million cell updates/sec

Title: US-10-681-199-1
Perfect score: 1263
Sequence: 1 atgctcttcaggtagcga.....gaacagaaactaaatcttaa 1263

Scoring table: OLIGO NUC
Gapop 60.0 , Gapext 60.0

Searched: 4390206 seqs, 2959870667 residues

Word size : 0

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : N_Geneseq_16Dec04.*
1: Geneseqn1980s.*
2: Geneseqn1990s.*
3: Geneseqn2000s.*
4: Geneseqn2001as.*
5: Geneseqn2001bs.*
6: Geneseqn2002as.*
7: Geneseqn2002bs.*
8: Geneseqn2003as.*
9: Geneseqn2003bs.*
10: Geneseqn2003cs.*
11: Geneseqn2003ds.*
12: Geneseqn2004as.*
13: Geneseqn2004bs.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	1263	100.0	1263	9 ADB16964	ADB16964 Human DYX
2	1263	100.0	1993	9 ADB16965	ADB16965 cDNA sequ
3	1049	83.1	1641	10 ADC30210	ADC30210 Human nov
4	1002	79.3	1263	9 ADB16939	ADB16939 Pygmy chi
5	998	79.0	1559	11 ADM01890	ADM01890 Human cDN
6	960	76.0	1263	9 ADB16933	ADB16933 Chimpanze
7	859	68.0	1263	9 ADB16935	ADB16935 Gorilla D
8	696	55.1	1263	9 ADB16937	ADB16937 Orangutan
9	524	41.5	608	10 ADC32116	ADC32116 Human nov
10	326	25.8	488	9 ACH35463	ACH35463 Human end
11	270	21.4	458	9 ACH23091	ACH23091 Human adu
12	197	15.6	1383	5 AAS70018	AAS70018 DNA encod
13	184	14.6	49806	9 ADB16927	ADB16927 Human DYX
14	155	12.3	49939	9 ADB16928	ADB16928 Human DYX
15	126	10.0	164	3 AAC30498	AAC30498 Human sec
16	109	8.6	313	3 AAC26799	AAC26799 Human sec
17	60	4.8	174	3 AAA45298	AAA45298 Human sec
18	56	4.4	174	3 AAA45298	AAA45298 Human sec
19	33	2.6	1697	9 ADB16924	ADB16924 cDNA sequ
20	25	2.0	25	9 ADB16932	ADB16932 Reverse R

c	21	25	2.0	25	9	ADB16931	ADB16931 Forward R
	22	22	1.7	1316	2	AAC07163	AAC07163 Human lun
	23	22	1.7	1316	3	AAC79075	AAC79075 Human lun
	24	22	1.7	1316	4	AAD23150	AAD23150 Human lun
	25	22	1.7	1316	10	ADD67087	ADD67087 Human lun
	26	22	1.7	1316	10	ADE87592	ADE87592 Human lun
	27	22	1.7	1431	4	ABL01953	ABL01953 Drosophil
	28	22	1.7	5746	4	ABL01952	ABL01952 Drosophil
c	29	21	1.7	579	4	AH09057	AH09057 Human cDN
	30	21	1.7	2310	12	ADI81619	ADI81619 C. elegan
	31	21	1.7	310268	13	ABD32548	ABD32548 Human can
	32	20	1.6	480	4	AAI86685	AAI86685 Human pol
	33	20	1.6	481	6	ABL83144	ABL83144 Human ova
c	34	20	1.6	1339	10	ADF82325	ADF82325 Leukaemia
	35	20	1.6	2967	13	ADR08160	ADR08160 Full leng
	36	20	1.6	3075	13	ADR08053	ADR08053 Full leng
	37	20	1.6	3143	13	ADR66793	ADR66793 Human pro
	38	20	1.6	3143	13	ADR65890	ADR65890 Human pro
	39	20	1.6	3183	13	ADR08303	ADR08303 Full leng
	40	20	1.6	8073	6	ABL32754	ABL32754 Human imm
	41	20	1.6	9110	6	ABL34429	ABL34429 Human imm
	42	20	1.6	48551	6	AAS20800	AAS20800 Clostridi
c	43	20	1.6	84248	6	ABQ99651	ABQ99651 Human MS4
	44	20	1.6	110000	12	ADQ97138_3	Continuation (4 of
	45	20	1.6	110469	12	ADQ97337	ADQ97337 Human can

ALIGNMENTS

RESULT 1
ADB16964
ID ADB16964 standard; cDNA; 1263 BP.
XX
AC ADB16964;
XX
DT 20-NOV-2003 (first entry)
XX
DE Human DYXC1 cDNA with single nucleotide polymorphisms.
XX
KW Gene; ss; human; DYXC1; dyslexia; neurological disorder;
KW Chromosome 15q21; reading disability; phonological processing;
KW rapid naming; verbal short-term memory; single nucleotide polymorphism;
KW SNP.
XX
OS Homo sapiens.
XX
FH Key
FT CDS
FT Location/Qualifiers
FT 1..1263
FT /tag= a
FT /product= "DYXC1 protein"
FT replace(4,t)
FT /tag= b
FT /standard_name= "Single nucleotide polymorphism"
FT replace(572,a)
FT /tag= c
FT /standard_name= "Single nucleotide polymorphism"
FT replace(1249,t)
FT /tag= d
FT /standard_name= "Single nucleotide polymorphism"
FT replace(1259,g)
FT /tag= e
FT /standard_name= "Single nucleotide polymorphism"
XX
WO2003068814-A1.
XX
PD 21-AUG-2003.
XX
PF 12-FEB-2003; 2003WO-FI000110.
XX
PR 12-FEB-2002; 2002US-0355782P.
XX
PA (LICN) LICENTIA LTD.

PN WO2003068814-A1.
XX 21-AUG-2003.
XX 12-FEB-2003; 2003WO-FI000110.
XX 12-FEB-2002; 2002US-0355782P.
XX (LION) LICENTIA LTD.
XX Kere J, Taipale M, Nopola-Hemmi J, Kaminen N;
XX WPI; 2003-646482/61.
XX P-P8DB; ADB16923.
XX New isolated, purified DYX1C1 nucleic acid for studying brain processes,
XX e.g. reading, phonological processing, rapid naming or verbal short-term
XX memory, or for diagnosing dyslexia or assessing the predisposition to
XX dyslexia.
XX Claim 10; Page 48-50; 135pp; English.
XX This invention relates to a novel isolated human gene DYX1C1 that is
XX functionally related to dyslexia, more particularly it describes single
XX nucleotide polymorphisms thought to predispose an individual in to
XX developing dyslexia. This is a neurological disorder with a genetic basis
XX (DYX1C1 has been isolated to chromosome 15q21), which manifests itself as
XX a specific reading disability. Specifically, DYX1C1 is can be useful in
XX study of brain processes such as reading, phonological processing, rapid
XX naming and verbal short-term memory. Accordingly, the present invention
XX describes methods and materials for analysing allelic variations in the
XX DYX1C1 gene, and also provides DYX1C1 as an antigen for the production of
XX antibodies used in the diagnosis of dyslexia. This polynucleotide is the
XX cDNA sequence of the wild type human DYX1C1 mRNA of the invention.
XX
XX SQ Sequence 1993 BP; 693 A; 368 C; 430 G; 502 T; 0 U; 0 Other;
Query Match 100.0%; Score 1263; DB 9; Length 1993;
Best Local Similarity 100.0%; Pred. No. 0;
Matches 1263; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
1 ATGCTCTTTCAGGTTAGCGATTACAGCTGGCGAGCAGAGACCTGCGGTCTTCTGCT 60
369 ATGCTCTTTCAGGTTAGCGATTACAGCTGGCGAGCAGAGACCTGCGGTCTTCTGCT 428
61 CTGCCCCCTCAAAGGGCGTGTGGCTCAGACACGACGAGCTGTTCTGACGGAACCTATCTG 120
429 CTGCCCCCTCAAAGGGCGTGTGGCTCAGACACGACGAGCTGTTCTGACGGAACCTATCTG 488
121 AAGGTCAACTTCTCTCCATTTTATTGAGGCAATTTCTTATGCTCCCATAGACGATGAG 180
489 AAGGTCAACTTCTCTCCATTTTATTGAGGCAATTTCTTATGCTCCCATAGACGATGAG 548
181 AGCAGCAAGCAAGATGGGAATGACACCATGCTCTTCACTTGTATATAAAGAGAGCG 240
549 AGCAGCAAGCAAGATGGGAATGACACCATGCTCTTCACTTGTATATAAAGAGAGCG 608
241 GCCATGTGGGAGACCTTTCTGTGACGGGTGTGACAAAGAGATGATGCAAGAAATAGA 300
609 GCCATGTGGGAGACCTTTCTGTGACGGGTGTGACAAAGAGATGATGCAAGAAATAGA 668
301 GAAATAATCTATTTTACAAGCAAGAGAGAGCAAGAAAGCTTACAGAGCAAAAGCTGCA 360
669 GAAATAATCTATTTTACAAGCAAGAGAGAGCAAGAAAGCTTACAGAGCAAAAGCTGCA 728
361 GCAGCGGGAGATCAAAATACGCACTAGTGTGATGATGATGATGATGATGATGATGATG 420
729 GCAGCGGGAGATCAAAATACGCACTAGTGTGATGATGATGATGATGATGATGATGATG 788
421 AGGAAAAAATAGAGATATGAAGAAATGAACGATATAAAGGCACTAAAGCAATTTGAA 480
789 AGGAAAAAATAGAGATATGAAGAAATGAACGATATAAAGGCACTAAAGCAATTTGAA 848

QY 481 GCCTGGAAGAAATATCAAGGAAGAAAGCTGAGGAGCAAAAGAAATTTCAAGAGAGAGAAA 540
DB 849 GCCTGGAAGAAATATCAAGGAAGAAAGCTGAGGAGCAAAAGAAATTTCAAGAGAGAGAAA 908
QY 541 TTATGTCACAAAAGAAAAGCAAAATTAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 600
DB 909 TTATGTCACAAAAGAAAAGCAAAATTAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 968
QY 601 ACTAGAAATTTGGCAATCTAGAAATCTTCTCTCAAAGGGAGAGAAATTTCAAGAGAGAG 660
DB 969 ACTAGAAATTTGGCAATCTAGAAATCTTCTCTCAAAGGGAGAGAGAAATTTCAAGAGAG 1028
QY 661 ACTGAGAGTTAAAGGAGAGACAGTATTTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 720
DB 1029 ACTGAGAGTTAAAGGAGAGACAGTATTTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 1088
QY 721 AACTTTACCCCTCGAGATTTCCCAACAGCTCTTCTGTAATCAACAAGTACAGAGAGAGAG 780
DB 1089 AACTTTACCCCTCGAGATTTCCCAACAGCTCTTCTGTAATCAACAAGTACAGAGAGAG 1148
QY 781 GAGTGGCTACACAAAGCTGAGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 840
DB 1149 GAGTGGCTACACAAAGCTGAGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 1208
QY 841 TGGCATTTAAAG 900
DB 1209 TGGCATTTAAAG 1268
QY 901 TTTCGACGAG 960
DB 1269 TTTCGACGAG 1328
QY 961 AATAAGATGCCATTTATTTGTAATTTGTAATTTGTAATTTGTAATTTGTAATTTGTAAT 1020
DB 1329 AATAAGATGCCATTTATTTGTAATTTGTAATTTGTAATTTGTAATTTGTAATTTGTAAT 1388
QY 1021 CACAGGCTATTGAGAGATTTCTTAAGGCACTGGAAATTTATGATGCCACCTGTTACAGAC 1080
DB 1389 CACAGGCTATTGAGAGATTTCTTAAGGCACTGGAAATTTATGATGCCACCTGTTACAGAC 1448
QY 1081 AATCTTAATGCAAGAAATGAAGGACATGTAGAGCTGGAACAGAGATTTCTGCACTAGAA 1140
DB 1449 AATCTTAATGCAAGAAATGAAGGACATGTAGAGCTGGAACAGAGATTTCTGCACTAGAA 1508
QY 1141 TTGTATGTAGAGAGCTTACAGGATTTATGAAGGCACTTAAAGATTTGATTCATCCAAAGAA 1200
DB 1509 TTGTATGTAGAGAGCTTACAGGATTTATGAAGGCACTTAAAGATTTGATTCATCCAAAGAA 1568
QY 1201 ATTGTACAAATTTGATGCTGAGAGATTTGCGGAATTTGATTTCAAGGAGAGAGAGAGAG 1260
DB 1569 ATTGTACAAATTTGATGCTGAGAGATTTGCGGAATTTGATTTCAAGGAGAGAGAGAGAG 1628
QY 1261 TAA 1263
DB 1629 TAA 1631
RESULT 3
ADC30210
ID ADC30210 standard; cDNA; 1641 BP.
XX
AC ADC30210;
XX
DT 18-DEC-2003 (first entry)
XX
DE Human novel cDNA sequence, SEQ ID NO:292.
XX
KW Human; diagnostic; drug screening; forensics; gene mapping;
KW biodiversity assessment; Parkinson's disease; Alzheimer's disease;
KW neurodegenerative diseases; anaemia; platelet disorder; wound; burns;
KW ulcers; osteoporosis; autoimmune disease; cancer;
KW molecular weight marker; food supplement; antiparkinsonian; nootropic;
KW neuroprotective; anti-nausea; anticoagulant; thrombolytic; vulnerary;

KW antiulcer; osteopathic; immunosuppressive; antiinflammatory; cytostatic;
KW gene therapy; chromosome 15q21.3; gene; ss.
OS Homo sapiens.
XX WO2003029271-A2.
XX 10-APR-2003.
XX 24-SEP-2002; 2002WO-US030474.
XX 24-SEP-2001; 2001US-0324631P.
XX (HYSE-) HYSEQ INC.
XX Tang TY, Zhang J, Ren F, Xue AJ, Zhao QA, Wang J, Wehrman T;
PI Zhou P, Ghosh M, Wang D, Ma Y, Asundi V, Wang Z, Weng G;
PI Haley-Vicente D, Drmanac RT;
XX WPI; 2003-371981/35.
DR P-PSDB; ADC31181.
XX New polynucleotide and polypeptide useful for diagnosing, preventing or
PT treating conditions such as neurodegenerative diseases, anemias, platelet
PT disorders, wounds, burns, ulcers, osteoporosis, autoimmune diseases or
PT cancer.
XX Claim 1; SEQ ID NO 292; 1185pp; English.
XX The invention relates to 971 novel human cDNA sequences (ADC29919-
CC ADC30889) and the polypeptides they encode (ADC30890-ADC31860). The
CC invention also relates to nucleic acid sequences over 99% identical with
CC the novel human cDNAs. The invention additionally encompasses expression
CC vectors and host cells comprising a nucleic acid of the invention; the
CC recombinant production of a polypeptide of the invention; an antibody
CC against a polypeptide of the invention; a method of detecting
CC polynucleotides or polypeptides of the invention; and methods of
CC identifying a compound which binds to a polypeptide of the invention. The
CC invention further discloses methods of preventing, treating or
CC ameliorating a medical condition; kits comprising polynucleotide probes
CC and/or monoclonal antibodies for carrying out the methods of the
CC invention; methods for the identification of compounds that modulate the
CC expression or activity of the polynucleotide and/or polypeptide; and 767
CC contig sequences corresponding to the cDNA sequences of the invention
CC (ADC31861-ADC32627) and the polypeptides encoded by the contigs (ADC32628
CC -ADC33394). The nucleic acids and polypeptides of the invention are
CC useful in diagnostics, drug screening, forensics, gene mapping, in the
CC identification of mutations responsible for genetic disorders or other
CC traits, for assessing biodiversity, and in producing many other types of
CC data and products dependent on DNA and amino acid sequences. They are
CC also used for treating diseases such as Parkinson's disease, Alzheimer's
CC disease and other neurodegenerative diseases, anaemia, platelet
CC disorders, wounds, ulcers, osteoporosis, autoimmune diseases or
CC cancer. The nucleic acids may also be used as hybridisation probes or
CC primers, and in the recombinant production of a protein. The polypeptides
CC are also useful in generating antibodies, as molecular weight markers,
CC and as food supplements. The present sequence represents a specifically
CC claimed human cDNA sequence of the invention. Note: The sequence data for
CC this patent did not form part of the printed specification, but was
CC obtained in electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences.
XX Sequence 1641 BP; 614 A; 272 C; 331 G; 424 T; 0 U; 0 Other;
S0 Query Match 83.1%; Score 1049; DB 10; Length 1641;
Best Local Similarity 100.0%; Pred. No. 0;
Matches 1049; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 ATGCTCTTCAAGTTAGCGATTACAGCTGGCAGACGACGACGACGCTTCTGTCT 60
DB 104 ATGCTCTTCAAGTTAGCGATTACAGCTGGCAGACGACGACGACGCTTCTGTCT 163
QY 61 CTGCCCTCAAAGCGTGTGCGTCAGACACGCGGCTGTTCTGCACGAAAACTATCTG 120

DB 164 CTGCCCTCAAAGCGTGTGCGTCAGACACGCGGCTGTTCTGCACGAAAACTATCTG 223
QY 121 AAGGTCAACTTCTCTCTCCATTTTATTTGAGGCATTTCTTTATGCTCCCATAGACGATGAG 180
DB 224 AAGGTCAACTTCTCTCTCCATTTTATTTGAGGCATTTCTTTATGCTCCCATAGACGATGAG 283
QY 181 AGCAGCAAAAGCAAAAGATTGGGAATGACACCATTTGTCTTCACTTTGTATATAAAAAAGAGCG 240
DB 284 AGCAGCAAAAGCAAAAGATTGGGAATGACACCATTTGTCTTCACTTTGTATATAAAAAAGAGCG 343
QY 241 GCCATGTGGGAGACCCCTTTCTGTGACGGGTGTTGACAAAGAGATGATGCAAAAGATTAGA 300
DB 344 GCCATGTGGGAGACCCCTTTCTGTGACGGGTGTTGACAAAGAGATGATGCAAAAGATTAGA 403
QY 301 GAAAAATCTATTTTACAGCACAAAGAGAGAGCAAAAGAGAGCTACAGAGCAAAAGCTGCA 360
DB 404 GAAAAATCTATTTTACAAAGCACAAAGAGAGAGCAAAAGAGAGCTACAGAGCAAAAGCTGCA 463
QY 361 GCAAGCGGGAAGATCAAAAATACGCACCTAAGTGTCTATGATGAAGATTGAAGAAGAGAGAG 420
DB 464 GCAAGCGGGAAGATCAAAAATACGCACCTAAGTGTCTATGATGAAGATTGAAGAAGAGAGAG 523
QY 421 AGGAAAAAATAGAGATATGAAAGAAAAATGAACGGATATAAAAGCCACTAAAGCATTTGAA 480
DB 524 AGGAAAAAATAGAGATATGAAAGAAAAATGAACGGATATAAAAGCCACTAAAGCATTTGAA 583
QY 481 GCCTGGAAGAATATCAAGAAAAAGCTGAGGAGCAAAAAAATTCAGAGAGAGAGAGAAA 540
DB 584 GCCTGGAAGAATATCAAGAAAAAGCTGAGGAGCAAAAAAATTCAGAGAGAGAGAGAAA 643
QY 541 TTATGTCAAAAAGAAAGCAAAATTTAAAGAGGAGAGAAAAAATAAATAATTAAGAGTCTT 600
DB 644 TTATGTCAAAAAGAAAGCAAAATTTAAAGAGGAGAGAAAAAATAAATAATTAAGAGTCTT 703
QY 601 ACTAGAAATTTGGCATCTAGAAATCTTGCTCCAAAAGGGAGAAATTCAGAAAAATATATTT 660
DB 704 ACTAGAAATTTGGCATCTAGAAATCTTGCTCCAAAAGGGAGAAATTCAGAAAAATATATTT 763
QY 661 ACTAGAGAGTTAAAGGAGAGACAGTATTCCTGCTCTCTGCTCTGTTGGCAGTATTAATAATC 720
DB 764 ACTAGAGAGTTAAAGGAGAGACAGTATTCCTGCTCTCTGCTCTGTTGGCAGTATTAATAATC 823
QY 721 AACTTTACCCCTCGAGTATTTCCCAACAGCTCTTCGTAATCAACAAGTAGCAGAGAGAGAG 780
DB 824 AACTTTACCCCTCGAGTATTTCCCAACAGCTCTTCGTAATCAACAAGTAGCAGAGAGAGAG 883
QY 781 GAGTGGCTACACAAACAAAGCTGAGGCAAGAGAGCAATGAATACTGACATAGCTGAACCTT 840
DB 884 GAGTGGCTACACAAACAAAGCTGAGGCAAGAGAGCAATGAATACTGACATAGCTGAACCTT 943
QY 841 TGGCATTTAAAG 900
DB 944 TGGCATTTAAAG 1003
QY 901 TTTGCAACGGAAGAACTATTTGGCAGCTATCAATGCATATTAATTTAGCCATTAAGACTAAAT 960
DB 1004 TTTGCAACGGAAGAACTATTTGGCAGCTATCAATGCATATTAATTTAGCCATTAAGACTAAAT 1063
QY 961 AATAAGATGACCATTTATTTGAAACCGGGCTGTTGCCACCTTAATAAATAAATAAATAA 1020
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QY 1021 CACAGGCTATTTGAAGATTCTTCTAAGGC 1049
DB 1124 CACAGGCTATTTGAAGATTCTTCTAAGGC 1152
RESULT 4
ADB16939
ID ADB16939 standard; cDNA; 1263 BP.
XX
AC ADB16939;

XX 20-NOV-2003 (first entry)
 XX Pygmy chimpanzee DYXC1 cDNA sequence.
 DE gene; ss: pygmy chimpanzee; DYXC1; dyslexia; neurological disorder;
 KW reading disability; phonological processing; rapid naming;
 KW verbal short-term memory.
 XX Pan paniscus.
 XX Key Location/Qualifiers
 FT 1..1263
 FT /*tag= a
 FT /product= "DYXC1 protein"
 XX WO2003068814-A1.
 XX 21-AUG-2003.
 XX 12-FEB-2003; 2003WO-FI000110.
 XX 12-FEB-2002; 2002US-0355782P.
 XX (LICN) LICENTIA LTD.
 XX Kere J, Taipale M, Nopola-Hemmi J, Kaminen N;
 PI WPI; 2003-646482/61.
 DR P-PSDB; ADB16940.
 XX New isolated, purified DYXC1 nucleic acid for studying brain processes,
 PT e.g. reading, phonological processing, rapid naming or verbal short-term
 PT memory, or for diagnosing dyslexia or assessing the predisposition to
 PT dyslexia.
 XX Claim 29; Page 126-128; 135pp; English.
 XX This invention relates to a novel isolated human gene DYXC1 that is
 CC functionally related to dyslexia, more particularly it describes single
 CC nucleotide polymorphisms thought to predispose an individual in to
 CC developing dyslexia. This is a neurological disorder with a genetic basis
 CC (DYXC1 has been isolated to chromosome 15q21), which manifests itself as
 CC a specific reading disability. Specifically, DYXC1 is can be useful in
 CC study of brain processes such as reading, phonological processing, rapid
 CC naming and verbal short-term memory. Accordingly, the present invention
 CC describes methods and materials for analysing allelic variations in the
 CC DYXC1 gene, and also provides DYXC1 as an antigen for the production of
 CC antibodies used in the diagnosis of dyslexia. This polynucleotide
 CC sequence is the pygmy chimpanzee DYXC1 cDNA homologous to the human DYXC1
 CC gene of the invention.
 XX Sequence 1263 BP; 493 A; 213 C; 274 G; 283 T; 0 U; 0 Other;
 SQ
 Query Match 79.3%; Score 1002; DB 9; Length 1263;
 Best Local Similarity 99.6%; Pred. No. 0;
 Matches 1252; Conservative 0; Mismatches 5; Indels 0; Gaps 0;
 QY 7 CTTCAAGGCTGTGGCTCAGACACGACGCTGTTCTGCACGGAACCTATCTGAAGGTC 126
 DB 7 CTTCAAGGCTGTGGCTCAGACACGACGCTGTTCTGCACGGAACCTATCTGAAGGTC 126
 QY 67 CTTCAAGGCTGTGGCTCAGACACGACGCTGTTCTGCACGGAACCTATCTGAAGGTC 126
 DB 67 CTTCAAGGCTGTGGCTCAGACACGACGCTGTTCTGCACGGAACCTATCTGAAGGTC 126
 QY 127 AACTTTCTCCATTTTATTTAGGCAATTTCTTTATGCTCCATAGACGATGAGGACG 186
 DB 127 AACTTTCTCCATTTTATTTAGGCAATTTCTTTATGCTCCATAGACGATGAGGACG 186
 QY 187 AAAGCAAGATTTGGGAATGACACCATTTGCTTACCTTGTATATAAAGAACGGCCATG 246
 DB 187 AAAGCAAGATTTGGGAATGACACCATTTGCTTACCTTGTATATAAAGAACGGCCATG 246

RESULT 5

ADM01890

ID ADM01890 standard; cDNA; 1559 BP.

QY 247 TGGGAGACCCCTTTCTGTGACGGGTGTTGACAAAGAGATGATGCCAAAGAAATTAGAGAAAAA 306
 DB 247 TGGGAGACCCCTTTCTGTGACGGGTGTTGACAAAGAGATGATGCCAAAGAAATTAGAGAAAAA 306
 QY 307 TCTATTTTACAGCAACAAGAGAGACCAAAAGAGCTTACAGAGCAAAAGCTCTCAGCAAG 366
 DB 307 TCTATTTTACAGCAACAAGAGAGACCAAAAGAGCTTACAGAGCAAAAGCTCTCAGCAAG 366
 QY 367 CGGGAAGATCAAAAATACGCACCTAAGTGTGATGATGAAGATTGAAGAAGAGAGGAAA 426
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 QY 427 AAAATAGAGATATGAAAGAAAATGAACGATAAAAGCCACTAAAGCATTGAGAGCTGG 486
 DB 427 AAAATAGAGATATGAAAGAAAATGAACGATAAAAGCCACTAAAGCATTGAGAGCTGG 486
 QY 487 AAAGAAATATCAAAAGAAAAGCTGAGGAGCAAAAAAATTCAGAGAGAGAGAAAATATCT 546
 DB 487 AAAGAAATATCAAAAGAAAAGCTGAGGAGCAAAAAAATTCAGAGAGAGAGAAAATATCT 546
 QY 547 CAAAAAGAAAAGCAAAATTAAGAGAGAGAAAATAAATAATAAGAGTCTTACTAGA 606
 DB 547 CAAAAAGAAAAGCAAAATTAAGAGAGAGAAAATAAATAATAAGAGTCTTACTAGA 606
 QY 607 AATTTGGCATCTAGAAATCTTGCTCCAAAAGGGAGAAAATTCAGAAAAATATATTACTGAG 666
 DB 607 AATTTGGCATCTAGAAATCTTGCTCCAAAAGGGAGAAAATTCAGAAAAATATATTACTGAG 666
 QY 667 AAGTTAAGGAGACAGATTTCTCTGCTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 726
 DB 667 AAGTTAAGGAGACAGATTTCTCTGCTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 726
 QY 727 ACCCTCTCGAGTATTCCTCAACAGCTCTTCTGCTGAATCAAGTAGCAGAGAGAGAGTGG 786
 DB 727 ACCCTCTCGAGTATTCCTCAACAGCTCTTCTGCTGAATCAAGTAGCAGAGAGAGAGTGG 786
 QY 787 CTACACAAACAGCTGAGGACGAGAGCAATGATGATGATGATGATGATGATGATGATGATGAT 846
 DB 787 CTGACACAAACAGCTGAGGACGAGAGCAATGATGATGATGATGATGATGATGATGATGAT 846
 QY 847 TTAAAG 906
 DB 847 TTAAAG 906
 QY 907 ACAGAAAACTATTTGGCAGCTATCAATGCATATAATTTAGCCATAGACTAAATAATAAG 966
 DB 907 ACAGAAAACTATTTGGCAGCTATCAATGCATATAATTTAGCCATAGACTAAATAATAAG 966
 QY 967 ATGCCACTATTGTATTGAAACGGGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 1026
 DB 967 ATGCCACTATTGTATTGAAACGGGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTG 1026
 QY 1027 GCTATTGAAGATTTCTTAAGGCACTGGAATTTATGATGATGATGATGATGATGATGATGAT 1086
 DB 1027 GCTATTGAAGATTTCTTAAGGCACTGGAATTTATGATGATGATGATGATGATGATGATGAT 1086
 QY 1087 AATGCAAGATGAGGACGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 1146
 DB 1087 AATGCAAGATGAGGACGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 1146
 QY 1147 GTAGAGAGCTTACAGGATTTGAGCGGCACTTAAAGATTGATGATGATGATGATGATGATGAT 1206
 DB 1147 GTAGAGAGCTTACAGGATTTGAGCGGCACTTAAAGATTGATGATGATGATGATGATGATGAT 1206
 QY 1207 CAAATTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 1263
 DB 1207 CAAATTGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGATGAT 1263

XX 21-AUG-2003.
PD 12-FEB-2003; 2003WO-FI000110.
XX PF
XX 12-FEB-2002; 2002US-0355782P.
PR
XX (LICN) LICENTIA LTD.
PA
XX Kere J, Taipale M, Nopola-Hemmi J, Kaminen N;
PI
XX WPI; 2003-646482/51.
DR P-PSDB; ADB16934.
XX
XX New isolated, purified DYXC1 nucleic acid for studying brain processes,
PT e.g. reading, phonological processing, rapid naming or verbal short-term
PT memory, or for diagnosing dyslexia or assessing the predisposition to
PT dyslexia.
XX
PS Claim 29; Page 116-118; 135pp; English.
XX
CC This invention relates to a novel isolated human gene DYXC1 that is
CC functionally related to dyslexia, more particularly it describes single
CC nucleotide polymorphisms thought to predispose an individual in to
CC developing dyslexia. This is a neurological disorder with a genetic basis
CC (DYXC1 has been isolated to chromosome 15q21), which manifests itself as
CC a specific reading disability. Specifically, DYXC1 is can be useful in
CC study of brain processes such as reading, phonological processing, rapid
CC naming and verbal short-term memory. Accordingly, the present invention
CC describes methods and materials for analysing allelic variations in the
CC DYXC1 gene, and also provides DYXC1 as an antigen for the production of
CC antibodies used in the diagnosis of dyslexia. This polynucleotide
CC sequence is the chimpanzee DYXC1 cDNA homologous to the human DYXC1 gene
CC of the invention.
XX
SQ Sequence 1263 BP; 494 A; 213 C; 272 G; 284 T; 0 U; 0 Other;
Query Match 76.0%; Score 960; DB 9; Length 1263;
Best Local Similarity 99.6%; Pred. No. 0;
Matches 1210; Conservative 0; Mismatches 5; Indels 0; Gaps 0;
49 GTCTTTCTGCTCTGCCCCCTCAAGCGTGTGCGTCAGAGCAGCGGTCTTCTGACG 108
49 GTCTTTCTGCTCTGCCCCCTCAAGCGTGTGCGTCAGAGCAGCGGTCTTCTGACG 108
109 GAAACTATCTGAAGTCAATCTTCTCCATTTTATTGAGGCAATTTCTTATGCTCCC 168
109 GAAACTATCTGAAGTCAATCTTCTCCATTTTATTGAGGCAATTTCTTATGCTCCC 168
169 ATAGACGATGAGAGCAGCAAGCAAGATTTGGGAATGACACCATTTGCTTACCTTGTAT 228
169 ATAGACGATGAGAGCAGCAAGCAAGATTTGGGAATGACACCATTTGCTTACCTTGTAT 228
229 AAAAAAGAGCGGCCATGTGGGAGACCTTTCTGTGACGGGTGTGACAAAGAGATGATG 288
229 AAAAAAGAGCGGCCATGTGGGAGACCTTTCTGTGACGGGTGTGACAAAGAGATGATG 288
289 CAAGAATTTAGAGAAAATCTATTTTACAGCACAAGAGAGAGCAAGAAAGAGCTACAGAA 348
289 CAAGAATTTAGAGAAAATCTATTTTACAGCACAAGAGAGAGCAAGAAAGAGCTACAGAA 348
349 GCAAAAGCTGAGCAAGCGGGAAGATCAAAAATACGCACTAAGTGTGATGATGAAGATT 408
349 GCAAAAGCTGAGCAAGCGGGAAGATCAAAAATACGCACTAAGTGTGATGATGAAGATT 408
409 GAAGAAGAGAGAGAGAAAATATAGAGATATGAAGAANAATGAACGGATTAAGCCACT 468
409 GAAGAAGAGAGAGAGAAAATATAGAGATATGAAGAANAATGAACGGATTAAGCCACT 468
469 AAAGCATTTGAGCGCTGGAAGAAATATCAAGAAAGCTGAGGACCAAAAAAATTCAG 528
469 AAAGAAATTTGGAAGCGCTGGAAGAAATATCAAGAAAGCTGAGGAGCAAAAAAATTCAG 528

QY 529 AGAGAGAGAGAAATTTATGTCAAAAAGAGAAAGCAAAATTTAAAGAGGAGAAATATAAA 588
DB |||||||||||||||||||||||||||||||||||||||||||||||||||||||||| 588
QY 589 TATAAGAGTCTTACTAGAAAATTTGSCATCTAGAAATCTTGTCCAAAAGGGAGAAATTC 648
DB |||||||||||||||||||||||||||||||||||||||||||||||||||||||||| 648
QY 649 GAAATATATATTTACTGAGAAGTTAAAGGAAGACAGTATTTCTCTGCTCTCTGTGGC 708
DB |||||||||||||||||||||||||||||||||||||||||||||||||||||||||| 708
QY 709 AGTATTAATCAACTTTTACCCCTCGAGTATTTCCCAACAGCTCTTCTGGAATCACAAGTA 768
DB |||||||||||||||||||||||||||||||||||||||||||||||||||||||||| 768
QY 769 GCAGAAGAGGAGGAGTGGCTACACAAACAACTGAGGACGAGGACGAGCAATGAATCTGAC 828
DB |||||||||||||||||||||||||||||||||||||||||||||||||||||||||| 828
QY 829 ATAGCTGAATCTTTCGATTTTAAAGAGAAAGAAAGAAACCCAGAAATGTTGAAGATAAA 888
DB |||||||||||||||||||||||||||||||||||||||||||||||||||||||||| 888
QY 889 GGAACAAATTTGTTGCAACGAAAACCTATTTGGCAGCTATCAATGCATATTAATTTAGCC 948
DB |||||||||||||||||||||||||||||||||||||||||||||||||||||||||| 948
QY 949 ATAGACTTAAATATAAGATCCACTATTTGTTTGAACCGGCTGCTTGCCACCTTAA 1008
DB |||||||||||||||||||||||||||||||||||||||||||||||||||||||||| 1008
QY 1009 CTAAAAAATTTACACAAGGCTTATGAAGATTTCTTCTAAGGCACTGGAATTTATTTGATGCA 1068
DB |||||||||||||||||||||||||||||||||||||||||||||||||||||||||| 1068
QY 1069 CCTGTTACAGACAAATGCTAATGCAAGAAATGAAGGACATGTACGAGTGGAAACAGCATTC 1128
DB |||||||||||||||||||||||||||||||||||||||||||||||||||||||||| 1128
QY 1129 TGTCAACTAGAAATTTGATGTAAGAGGCTTACAGGATTTATGAAGCGGCACTTAAGATTGAT 1188
DB |||||||||||||||||||||||||||||||||||||||||||||||||||||||||| 1188
QY 1189 CCATCCACAAAATTTGTACAAAATTTGATCTGAGAAGATTTGGAATTTAAATTCAGGAACA 1248
DB |||||||||||||||||||||||||||||||||||||||||||||||||||||||||| 1248
QY 1249 GAACTAAATCTTAA 1263
DB |||||||||||||||||||||||||||||||||||||||||||||||||||||||||| 1263
RESULT 7
ADBI6935
ID ADBI6935 standard; cDNA; 1263 BP.
XX AC ADBI6935;
XX XX
DT 20-NOV-2003 (first entry)
XX
DE Gorilla DYXC1 cDNA sequence.
XX
KW Gene; ss; gorilla; DYXC1; dyslexia; neurological disorder;
KW reading disability; phonological processing; rapid naming;
XX verbal short-term memory.
OS Gorilla gorilla.
XX
FH Key Location/Qualifiers
FT CDS 1..1263
FT /*tag= a
FT /product= "DYXC1 protein"

CC 38043 cDNA sequences, appearing as ACH12789-ACH50831, whose sequence was
CC determined by the technique of SBH (sequencing by hybridisation). Also
CC included is a purified polypeptide comprising a sequence corresponding to
CC a reading frame of the novel polynucleotide. The nucleic acid sequences
CC are useful in diagnostics as expressed sequence tags (EST) for
CC identifying expressed genes or for physical mapping of the human genome,
CC in forensics, in assessing biodiversity, or in identifying mutations
CC responsible for genetic disorders and other traits. The nucleotide
CC sequences are also useful as hybridisation probes, as oligomers for PCR,
CC for chromosome and gene mapping, in the recombinant production of
CC protein, or in generating antisense DNA or RNA. The purified polypeptide
CC is useful for generating antibodies specific for it. The present sequence
CC is one of the 38043 isolated cDNA/EST sequences. Note: The sequence data
CC for this patent did not form part of the printed specification, but was
CC obtained in electronic format directly from USPTO at
CC seqdata.uspto.gov/sequence.html?DocID=20030073623
XX

SQ Sequence 488 BP; 141 A; 105 C; 126 G; 116 T; 0 U; 0 Other;

Query Match 25.8%; Score 326; DB 9; Length 488;

Best Local Similarity 100.0%; Pred. No. 4.1e-145;

Matches 326; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 80 GGGTCAGAGACGCGAGGTTCTGCACGAAACCTATCTGAAGTCAACTTCTCTCCAT 139

DB 73 GGGTCAGAGACGCGAGGTTCTGCACGAAACCTATCTGAAGTCAACTTCTCTCCAT 132

QY 140 TTTTATTTGAGGCATTTCTTTATGCTCCATAGACGATGAGCAGCAAGCAAGATTG 199

DB 133 TTTTATTTGAGGCATTTCTTTATGCTCCATAGACGATGAGCAGCAAGCAAGATTG 192

QY 200 GGAATGACACCATTTCTTCACTTGTATATAAAAGAGCGGCCCATGTGGAGACCCCTT 259

DB 193 GGAATGACACCATTTCTTCACTTGTATATAAAAGAGCGGCCCATGTGGAGACCCCTT 252

QY 260 CTGTGACGGGTGTTGACAAAGAGATGATGCAAGAGATTTAGAAAATCTATTTTACAG 319

DB 253 CTGTGACGGGTGTTGACAAAGAGATGATGCAAGAGATTTAGAAAATCTATTTTACAG 312

QY 320 CACAAGAGAGAGCAAAAGAGCTACAGAGCAAAAGCTGCAAGCGGGAAGATCAAA 379

DB 313 CACAAGAGAGAGCAAAAGAGCTACAGAGCAAAAGCTGCAAGCGGGAAGATCAAA 372

QY 380 AATACGCACTAAGTGTCTATGATGAAG 405

DB 373 AATACGCACTAAGTGTCTATGATGAAG 398.

RESULT 11

ID ACH23091

ACH23091 standard; cDNA; 458 BP.

AC ACH23091;

XX

XX 13-OCT-2003 (first entry)

DT Human adult ovary cDNA #1471.

DE Human; ss; sequencing by hybridisation; SBH; expressed sequence tag; EST;

XX genome mapping; biodiversity; genetic disorder.

XX Homo sapiens.

XX US2003073623-A1.

XX 17-APR-2003.

XX 30-JUL-2001; 2001US-00918995.

XX 30-JUL-2001; 2001US-00918995.

XX (DRNA/) DRMANAC R T.

XX (LABA/) LABAT I.

PA (STAC/) STACHE-CRAIN B.

PA (DICK/) DICKSON M C.

XX (JONE/) JONES L W.

PI Drmanac RT, Labat I, Stache-Crain B, Dickson MC, Jones LW;

XX WPI; 2003-615964/58.

XX New polynucleotide sequences obtained from various cDNA libraries, useful
PT as hybridization probes, as oligomers for PCR, for chromosome and gene
PT mapping, in the recombinant production of protein, or in generating
PT antisense DNA or RNA.

XX Claim 1; SEQ ID NO 10303; 44pp; English.

CC The invention relates to an isolated polynucleotide comprising any one of
CC 38043 cDNA sequences, appearing as ACH12789-ACH50831, whose sequence was
CC determined by the technique of SBH (sequencing by hybridisation). Also
CC included is a purified polypeptide comprising a sequence corresponding to
CC a reading frame of the novel polynucleotide. The nucleic acid sequences
CC are useful in diagnostics as expressed sequence tags (EST) for
CC identifying expressed genes or for physical mapping of the human genome,
CC in forensics, in assessing biodiversity, or in identifying mutations
CC responsible for genetic disorders and other traits. The nucleotide
CC sequences are also useful as hybridisation probes, as oligomers for PCR,
CC for chromosome and gene mapping, in the recombinant production of
CC protein, or in generating antisense DNA or RNA. The purified polypeptide
CC is useful for generating antibodies specific for it. The present sequence
CC is one of the 38043 isolated cDNA/EST sequences. Note: The sequence data
CC for this patent did not form part of the printed specification, but was
CC obtained in electronic format directly from USPTO at
CC seqdata.uspto.gov/sequence.html?DocID=20030073623
XX

SQ Sequence 458 BP; 170 A; 86 C; 97 G; 102 T; 0 U; 3 Other;

Query Match 21.4%; Score 270; DB 9; Length 458;

Best Local Similarity 100.0%; Pred. No. 2.3e-118;

Matches 270; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 773 AAGAGGAGGAGTGCTACACAAAGCTGAGGCGACGAGAGCAATGATCTGACATAG 832

DB 189 AAGAGGAGGAGTGCTACACAAAGCTGAGGCGACGAGAGCAATGATCTGACATAG 248

QY 833 CTGAACCTTTGCGATTTAAAGAGAGAGAAAGACCCAGATGTTGAAGGATAAGGAA 892

DB 249 CTGAACCTTTGCGATTTAAAGAGAGAGAAAGACCCAGATGTTGAAGGATAAGGAA 308

QY 893 ACAATTTGTTGCAACGGAACCTATTTGGCAGCTATCAATGCATATATTTAGCCATAA 952

DB 309 ACAATTTGTTGCAACGGAACCTATTTGGCAGCTATCAATGCATATATTTAGCCATAA 368

QY 953 GACTAAATATAAGATGCCACTATTGTATTTGAAACCGGGCTTGCACCTATAAACTAA 1012

DB 369 GACTAAATATAAGATGCCACTATTGTATTTGAAACCGGGCTTGCACCTATAAACTAA 428

QY 1013 AAAAATTACACAGGCTATTGAAGATCTT 1042

DB 429 AAAAATTACACAGGCTATTGAAGATCTT 458

RESULT 12

AAAS70018

ID AAAS70018 standard; cDNA; 1383 BP.

XX AAAS70018;

XX 13-FEB-2002 (first entry)

DT DNA encoding novel human diagnostic protein #5822.

XX Human; chromosome mapping; gene mapping; gene therapy; forensic;
XX food supplement; medical imaging; diagnostic; genetic disorder; ss.

OS Homo sapiens.
 XX WO200175067-A2.
 XX 11-OCT-2001.
 XX 30-MAR-2001; 2001WO-US008631.
 XX 31-MAR-2000; 2000US-00540217.
 PR 23-AUG-2000; 2000US-00649167.
 XX (HYSE-) HYSEQ INC.
 XX Dmanac RT, Liu C, Tang YT;
 XX WPI; 2001-639362/73.
 DR P-PSDB; ABG05831.
 XX New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity.
 XX Claim 1; SEQ ID NO 5822; 103pp; English.
 XX The invention relates to isolated polynucleotide (I) and polypeptide (II)
 CC sequences. (I) is useful as hybridisation probes, polymerase chain
 CC reaction (PCR) primers, oligomers, and for chromosome and gene mapping,
 CC and in recombinant production of (II). The polynucleotides are also used
 CC in diagnostics as expressed sequence tags for identifying expressed
 CC genes. (I) is useful in gene therapy techniques to restore normal
 CC activity of (II) or to treat disease states involving (II). (II) is
 CC useful for generating antibodies against it, detecting or quantitating a
 CC polypeptide in tissue, as molecular weight markers and as a food
 CC supplement. (II) and its binding partners are useful in medical imaging
 CC of sites expressing (II). (I) and (II) are useful for treating disorders
 CC involving aberrant protein expression or biological activity. The
 CC polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human diagnostic
 CC coding sequences of the invention. Note: The sequence data for this
 CC patent did not appear in the printed specification, but was obtained in
 CC electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX Sequence 1383 BP; 399 A; 343 C; 337 G; 304 T; 0 U; 0 Other;
 SQ Query Match 15.6%; Score 197; DB 5; Length 1383;
 Best Local Similarity 100.0%; Pred. No. 1.8e-83;
 Matches 197; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 172 GACGATGAGCAGCAAGCAAGATGGGAATGACACCATGTCTTCACCTTGTATAAA 231
 DB 709 GACGATGAGCAGCAAGCAAGATGGGAATGACACCATGTCTTCACCTTGTATAAA 768
 OY 232 AAAGAAGCGGCATGTGGGAGACCCCTTCTGTGACGGGTGTGTGCAAGAGATGATGCAA 291
 DB 769 AAAGAAGCGGCATGTGGGAGACCCCTTCTGTGACGGGTGTGTGCAAGAGATGATGCAA 828
 OY 292 AGAATTAGAGAAAATCTATTTCACAGCAAGAGAGAGCAAGAGAGCTTACAGAGCA 351
 DB 829 AGAATTAGAGAAAATCTATTTCACAGCAAGAGAGAGCAAGAGAGCTTACAGAGCA 888
 OY 352 AAAGCTGCAGCAAGCG 368
 DB 889 AAAGCTGCAGCAAGCG 905
 RESULT 13
 ADB16927
 ID ADB16927 standard; DNA; 49806 BP.

XX ADB16927;
 XX 20-NOV-2003 (first entry)
 XX Human DYXC1 DNA, chromosomal gene region nucleotides 50001-100000.
 DE ds; human; DYXC1; dyslexia; neurological disorder; chromosome 15q21;
 KW reading disability; phonological processing; rapid naming;
 KW verbal short-term memory.
 XX Homo sapiens.
 XX WO2003068814-A1.
 XX 21-AUG-2003.
 XX 12-FEB-2003; 2003WO-FI000110.
 PR 12-FEB-2002; 2002US-0355782P.
 XX (LICN) LICENTIA LTD.
 XX Kere J, Taipale M, Nopola-Hemmi J, Kaminen N;
 XX WPI; 2003-646482/61.
 XX New isolated, purified DYXC1 nucleic acid for studying brain processes,
 PT e.g. reading, phonological processing, rapid naming or verbal short-term
 PT memory, or for diagnosing dyslexia or assessing the predisposition to
 PT dyslexia.
 XX Claim 27; Page 69-83; 135pp; English.
 XX This invention relates to a novel isolated human gene DYXC1 that is
 CC functionally related to dyslexia, more particularly it describes single
 CC nucleotide polymorphisms thought to predispose an individual in to
 CC developing dyslexia. This is a neurological disorder with a genetic basis
 CC (DYXC1) has been isolated to chromosome 15q21, which manifests itself as
 CC a specific reading disability. Specifically, DYXC1 is can be useful in
 CC study of brain processes such as reading, phonological processing, rapid
 CC naming and verbal short-term memory. Accordingly, the present invention
 CC describes methods and materials for analysing allelic variations in the
 CC DYXC1 gene, and also provides DYXC1 as an antigen for the production of
 CC antibodies used in the diagnosis of dyslexia. This polynucleotide is the
 CC partial genomic sequence of the human DYXC1 chromosomal region
 CC (nucleotides 50001-100000) of the invention.
 XX Sequence 49806 BP; 12975 A; 10105 C; 10577 G; 15977 T; 0 U; 172 Other;
 SQ Query Match 14.6%; Score 184; DB 9; Length 49806;
 Best Local Similarity 99.6%; Pred. No. 2.8e-77;
 Matches 234; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 OY 404 AGATTGAAGAAGAGAGAGAGGAAAGAAATAGAAAGATATGAAGAAATGAACGGATAAAG 463
 DB 32444 AGATTGAAGAAGAGAGAGGAGGAAAGAAATAGAGATATGAAGAAATGAACGGATAAAG 32503
 OY 464 CCACATAAGCATTTGGAGCCCTGGAAAGAAATATCAAGAAAGCTGAGAGCAAAAAAAA 523
 DB 32504 CCACATAAGCATTTGGAGCCCTGGAAAGAAATATCAAGAAAGCTGAGAGCAAAAAAAA 32563
 OY 524 TTTCAGAGAGAGAGAAATTTGTCAAAAAGAAAGCAAAATTAAGAAAGAGGAGAAAAAAA 583
 DB 32564 TTTCAGAGAGAGAGAAATTTGTCAAAAAGAAAGCAAAATTAAGAAAGAGGAGAAAAAAA 32623
 OY 584 TAAATATAGAGTCTTACTAGAAATTTGGCATCTAGAAATCTTGCTCCAAAAG 638
 DB 32624 TAAATATAGAGTCTTACTAGAAATTTGGCATCTAGAAATCTTGCTCCAAAAG 32678
 RESULT 14
 ADB16928

ID ADB16928 standard; DNA; 49939 BP.
AC ADB16928;
XX
XX
XX 20-NOV-2003 (first entry)
DT
DE Human DYXC1 DNA, chromosomal gene region nucleotides 100001-150000.
DE
XX ds; human; DYXC1; dyslexia; neurological disorder; chromosome 15q21;
KW reading disability; phonological processing; rapid naming;
KW Verbal short-term memory.
XX
XX Homo sapiens.
OS
XX WO2003068814-A1.
PN
XX
XX 21-AUG-2003.
PD
XX 12-FEB-2003; 2003WO-FI000110.
PF
XX 12-FEB-2002; 2002US-0355782P.
PR
XX (LICN) LICENTIA LTD.
PA
XX Kere J, Taipale M, Nopola-Hemmi J, Kaminen N;
PI
XX WPI; 2003-646482/61.
XX
XX New isolated, purified DYXC1 nucleic acid for studying brain processes.
PT e.g. reading, phonological processing, rapid naming or verbal short-term
PT memory, or for diagnosing dyslexia or assessing the predisposition to
PT dyslexia.
XX
XX Claim 27; Page 83-97; 135pp; English.
XX
XX This invention relates to a novel isolated human gene DYXC1 that is
CC functionally related to dyslexia, more particularly it describes single
CC nucleotide polymorphisms thought to predispose an individual in to
CC developing dyslexia. This is a neurological disorder with a genetic basis
CC (DYXC1 has been isolated to chromosome 15q21), which manifests itself as
CC a specific reading disability. Specifically, DYXC1 is can be useful in
CC study of brain processes such as reading, phonological processing, rapid
CC naming and verbal short-term memory. Accordingly, the present invention
CC describes methods and materials for analysing allelic variations in the
CC DYXC1 gene, and also provides DYXC1 as an antigen for the production of
CC antibodies used in the diagnosis of dyslexia. This polynucleotide is the
CC partial genomic sequence of the human DYXC1 chromosomal region
CC (nucleotides 100001-150000) of the invention.
XX
XX Sequence 49939 BP; 13786 A; 10910 C; 11127 G; 14116 T; 0 U; 0 Other;
SQ
Query Match 12.3%; Score 155; DB 9; Length 49939;
Best Local Similarity 100.0%; Pred. No. 2e-63;
Matches 155; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 894 CAATTTGTTGCAACGGAACACTATTGGCAGCTATCAATGATCATATAATTAGCCATAAG 953
Db 14682 CAATTTGTTGCAACGGAACACTATTGGCAGCTATCAATGATCATATAATTAGCCATAAG 14741
QY 954 ACTAATATATAGATGCCACTATTGTTATTTGACCGGGCTGCTGCCACCTAAACTAAA 1013
Db 14742 ACTAATATATAGATGCCACTATTGTTATTTGACCGGGCTGCTGCCACCTAAACTAAA 14801
QY 1014 AAACCTTACACAGGCTATTGAAGATTCTTCTAAGG 1048
Db 14802 AAACCTTACACAGGCTATTGAAGATTCTTCTAAGG 14836
102(6)
RESULT 15
AAC30498
ID AAC30498 standard; CDNA; 164 BP.
XX
AC AAC30498;

XX 06-OCT-2000 (first entry)
DT
XX Human secreted protein 5' EST, SEQ ID NO: 34573.
DE
XX Human; 5' EST; expressed sequence tag; secreted protein; cDNA isolation;
KW gene therapy; chromosome mapping; ss.
XX
XX Homo sapiens.
OS
XX EP1033401-A2.
PN
XX 06-SEP-2000.
PD
XX 21-FEB-2000; 2000EP-00200610.
PF
XX 26-FEB-1999; 99US-0122487P.
PR
XX (GEST) GENSET.
PA
XX Dumas Milne Edwards J, Duclert A, Giordano J;
PI
XX WPI; 2000-500381/45.
XX
XX New nucleic acid that is a 5' expressed sequence tag (5' EST) for
PT obtaining cDNAs and genomic DNAs that correspond to 5' ESTs and for
PT diagnostic, forensic, gene therapy and chromosome mapping procedures.
PT
XX
XX Claim 1; SEQ ID NO 34573; 71pp + Sequence Listing; English.
XX
XX The present sequence is one of a large number of 5' ESTs derived from
CC mRNAs encoding secreted proteins. NO ORF has yet been conclusively
CC identified within the present sequence. The 5' ESTs were prepared from
CC total human RNAs or polyA+ RNAs derived from 30 different tissues. EST
CC sequences usually correspond mainly to the 3' untranslated region (UTR)
CC of the mRNA because they are often obtained from oligo-dT primed cDNA
CC libraries. Such ESTs are not well suited for isolating cDNA sequences
CC derived from the 5' ends of mRNAs and even in those cases where longer
CC cDNA sequences have been obtained, the full 5' UTR is rarely included. 5'
CC ESTs are derived from mRNAs with intact 5' ends and can therefore be used
CC to obtain full length cDNAs and genomic DNAs. 5' ESTs are also used in
CC diagnostic, forensic, gene therapy and chromosome mapping procedures.
CC They are used to obtain upstream regulatory sequences and to design
CC expression and secretion vectors
XX
XX Sequence 164 BP; 34 A; 43 C; 38 G; 47 T; 0 U; 2 Other;
SQ
Query Match 10.0%; Score 126; DB 3; Length 164;
Best Local Similarity 100.0%; Pred. No. 1.5e-49;
Matches 126; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 5 CTCCTTCAGGTTAGCGATTACAGCTGGCAGCAGACGAACTGCGGTCTTTCTGCTCTGCG 64
Db 1 CTCCTTCAGGTTAGCGATTACAGCTGGCAGCAGACGAACTGCGGTCTTTCTGCTCTGCG 60
QY 65 CCCTCAAAGGCGGTGCGTTCAGAGACACGACGCGGTGTTCTGCACGAAACTATCTGAAGG 124
Db 61 CCCTCAAAGGCGGTGCGTTCAGAGACACGACGCGGTGTTCTGCACGAAACTATCTGAAGG 120
QY 125 TCAACT 130
Db 121 TCAACT 126
Search completed: July 8, 2005, 07:48:42
Job time : 763 secs

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OM nucleic - nucleic search, using sw model

Run on: July 8, 2005, 04:50:29 ; Search time 5660 Seconds
(without alignments)
10812.533 Million cell updates/sec

Title: US-10-681-199-1
Perfect score: 1263
Sequence: 1 atgcctcttcaggtagcga.....gaacagaactaaatcttaa 1263

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 4708233 seqs, 24227607955 residues

Word size : 0

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : GenEmbl.*

1: gb_ba.*
2: gb_hg.*
3: gb_in.*
4: gb_om.*
5: gb_ov.*
6: gb_pat.*
7: gb_ph.*
8: gb_pl.*
9: gb_pr.*
10: gb_ro.*
11: gb_sts.*
12: gb_sy.*
13: gb_un.*
14: gb_vl.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1263	100.0	1993	9 AF337549	Homo sapi
2	998	79.0	1559	6 AX833451	Sequence
3	998	79.0	1559	9 AK095201	Sequence
4	997	78.9	1468	9 BC062564	Homo sapi
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22	138	10.9	153	9	AY17859287	Pan panis
23	138	10.9	153	9	AY17860187	Gorilla g
24	138	10.9	153	9	AY17861087	Pongo pyg
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ALIGNMENTS

RESULT 1
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LOCUS Homo sapiens EKNI (EKNI) mRNA, complete cds.
DEFINITION AF337549
ACCESSION AF337549
VERSION AF337549.1 GI:18478647
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE
AUTHORS
Taipale, M., Kaminen, N., Nopola-Hemmi, J., Haltia, T., Myllyluoma, B.,
Lytinen, H., Muller, K., Kaaranen, M., Lindsberg, P.J.,
Hannula-Jouppi, K. and Kere, J.

TITLE
A candidate gene for developmental dyslexia encodes a nuclear
tetrapeptide repeat domain dynamically regulated in
brain

JOURNAL
MEDLINE
PROC. NATL. ACAD. SCI. U.S.A. 100 (20), 11553-11558 (2003)
PUBMED
22882828
12954984

REFERENCE
AUTHORS
Taipale, M. and Kere, J.
TITLE
A gene disrupted by translocation breakpoint in chromosome 15q21
Unpublished

REFERENCE
AUTHORS
Taipale, M. and Kere, J.
TITLE
Direct Submission
JOURNAL
Submitted (17-JAN-2001) Finnish Genome Center, University of
Helsinki, Tukholmankatu 2, Helsinki 00014, Finland

FEATURES
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Qy 241 GCCATGTGGAGACCTTTCTGTGACGGGTCTGCACAAAGAGATGATGCAAGAAAT 300
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Db 1629 TAA 1631

RESULT 2
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LOCUS AX833451
DEFINITION Sequence 575 from Patent EP1347046.
ACCESSION AX833451
VERSION AX833451.1 GI:39919586
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Isegai, T., Sugiyama, T., Otsuki, T., Wakamatsu, A., Sato, H., Ishii, S.,
Yamamoto, J. I., Isono, Y., Hio, Y., Otsuka, K., Nagai, K., Irie, R.,
Tamechika, I., Seki, N., Yoshikawa, T., Otsuka, M., Nagahari, K. and
Masuho, Y.
TITLE Full-length cDNA sequences
JOURNAL Patent: EP 1347046-A 575 24-SEP-2003;
Research Association for Biotechnology (JRP)
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Query Match 79.0%; Score 998; DB 6; Length 1559;
Best Local Similarity 99.9%; Pred. No. 0;
Matches 1048; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ATGCCTCTTTCAGGTTAGCGATTACAGCTGGCAGCAGAGAGACTGCGGTCTTCTGCT 60
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 DEFINITION
 ACCESSION AK095201
 VERSION AK095201.1 GI:21754405
 KEYWORDS oligo capping; fis (full insert sequence).
 SOURCE Homo sapiens (human)

ORGANISM
REFERENCE
AUTHORS

Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1
 Ota, T., Suzuki, Y., Nishikawa, T., Otsuki, T., Sugiyama, T., Irie, R.,
 Wakamatsu, A., Hayaashi, K., Sato, H., Nagai, K., Kimura, K., Makita, H.,
 Sekine, M., Ohayashi, M., Nishi, T., Shibahara, T., Tanaka, T.,
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 Nagahori, K., Murakami, K., Yasuda, T., Iwayanagi, T., Wagatsuma, M.,
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 Sugawara, M., Takahashi, M., Kanda, K., Kamihara, K., Yokoi, T., Furuya, T.,
 Kikkawa, E., Omura, Y., Abe, K., Kimihara, K., Katauchi, N., Sato, K.,
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 Kumagai, A., Itakura, S., Fukuzumi, Y., Fujimori, Y., Komiyama, M.,
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 Nakajima, Y., Mizuno, T., Morinaga, M., Sasaki, M., Togashi, T.,
 Noguchi, T., Shirai, Y., Takahashi, Y., Nakagawa, K., Okumura, K.,
 Oyama, M., Hata, H., Watanabe, M., Komatsu, T., Mizushima-Sugano, J.,
 Nagase, T., Nomura, N., Kikuchi, H., Masuho, Y., Yamashita, R.,
 Nakai, K., Yada, T., Nakamura, Y., Ohara, O., Isogai, T. and Sugano, S.
 Complete sequencing and characterization of 21,243 full-length
 human cDNAs
 Nat. Genet. 36 (1), 40-45 (2004)
 14702039

TITLE
JOURNAL
PUBMED
REFERENCE
AUTHORS

2
 Kawakami, B., Sugiyama, A., Takemoto, M., Sugiyama, T., Irie, R.,
 Otsuki, T., Sato, H., Wakamatsu, A., Ishii, S., Yamamoto, J., Isono, Y.,
 Kawai-Hio, Y., Saito, K., Nishikawa, T., Kimura, K., Yamashita, H.,
 Matsuo, K., Nakamura, Y., Sekine, M., Kikuchi, H., Kanda, K.,
 Wagatsuma, M., Murakawa, K., Kanehori, K., Takahashi-Fujii, A.,
 Oshima, A., Suzuki, Y., Sugano, S., Nagahori, K., Masuho, Y., Nagai, K.
 and Isogai, T.
 NEDO human cDNA sequencing project
 Unpublished
 3 (bases 1 to 1559)
 Isogai, T. and Yamamoto, J.
 Direct Submission
 Submitted (04-JUL-2002) Takao Isogai, FLJ Project (HRI Team); 2-6-7
 Kazusa-Kamatari, Kisarazu, Chiba 292-0812, Japan
 (E-mail: genomics@hri.co.jp, Tel: 81-438-52-3975, Fax: 81-438-52-3986)

TITLE
JOURNAL
AUTHORS
TITLE
JOURNAL

3
 NEDO human cDNA sequencing project supported by Ministry of
 Economy, Trade and Industry of Japan; cDNA full insert sequencing:
 Research Association for Biotechnology (RAB); cDNA library
 construction: Helix Research Institute (HRI) (supported by Japan
 Key Technology Center etc.); 5'- & 3'-end one pass sequencing: RAB,
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 Evaluation; clone selection for full insert sequencing: HRI and
 RAB; annotation: HRI and RAB.
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LOCUS AX970431 489 bp DNA linear PAT 15-JAN-2004
DEFINITION Sequence 1234 from Patent EP1104808.
ACCESSION AX970431
VERSION AX970431.1 GI:40977781
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Dumas Milne Edwards,J.B., Jobert,S. and Giordano,J.Y.
TITLE ESTs and encoded human proteins
JOURNAL Patent: EP 1104808-A 1234 06-JUN-2001;
Genset (FR)
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DEFINITION EST and encoded human protein.
ACCESSION BD109150
VERSION BD109150.1 GI:23203968
KEYWORDS JP 2002010789-A/1227.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Edwards,J.B.D.M., Jobert,S. and Giordano,J.E.
TITLE EST and encoded human protein
JOURNAL Patent: JP 2002010789-A 1227 15-JAN-2002;
GENSET CORP
OS Homo sapiens (human)
PN JP 2002010789-A/1227
PD 15-JAN-2002
PF 07-AUG-2000 JP 2000280989
PR 05-AUG-1999 US 60/147499
PI JEAN BAPTISTE DUMAS MILNE EDWARDS,SEVELIN JOBERT,JEAN EVE PI
GIORDANO
PC C12N15/09, C12N15/09, C07K14/47, C07K16/18, C12N1/15, C12N1/19, PC
C12N1/21,
PC C12N5/10, C12P21/02, C12P21/08, C12Q1/68, C12N15/00, C12N5/00, PC
C12N15/00
CC EST and encoded human protein
FH Key Location/Qualifiers
FT CDS
57..488.
Location/Qualifiers
1..489
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
ORIGIN
Query Match 32.2%; Score 407; DB 6; Length 489;
Best Local Similarity 100.0%; Pred. No. 1.2e-196;
Matches 407; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 27 CTGGCAGCAGACGAGACTCGGGTCTTTCTGTCTCTGCCCCCTCAAAGCGGTGCGTCAG 86
Db 83 CTGGCAGCAGACGAGACTCGGGTCTTTCTGTCTCTGCCCCCTCAAAGCGGTGCGTCAG 142
QY 87 AGACACGACGCTGTTCTGCAGCGAAACTATCTGAAGTCAACTTTCCTCCATTTTATT 146
Db 143 AGACACGACGCTGTTCTGCAGCGAAACTATCTGAAGTCAACTTTCCTCCATTTTATT 202
QY 147 TGAGGCATTTCTTTATGTCCTCCATAGACGATGAGACGAGCAAAAGATTGGGAATGA 206
Db 203 TGAGGCATTTCTTTATGTCCTCCATAGACGATGAGACGAGCAAAAGATTGGGAATGA 262
QY 207 CACCATTTGTCCTTACCTTGTATATAAAGAGCGGCCCATGTGGAGACCCCTTTCTGTGAC 266
Db 263 CACCATTTGTCCTTACCTTGTATATAAAGAGCGGCCCATGTGGAGACCCCTTTCTGTGAC 322
QY 267 GGGTGTTCACAAAGAGATGATGCAAGAATTTAGAGAAAAATCTATTTTACAAGCACAGA 326
Db 323 GGGTGTTCACAAAGAGATGATGCAAGAATTTAGAGAAAAATCTATTTTACAAGCACAGA 382
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Db      229  AGCAGCAAAAGATTGGGAATGACACCATTTG 262
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AC021668      43886 bp      DNA      linear      HTG 29-MAR-2001
Homo sapiens chromosome 15 clone RP11-13306 map 15, LOW-PASS
SEQUENCE SAMPLING.
AC021668      4  GI:13488020
HTG; HTGS PHASE0.
SOURCE      Homo sapiens (human)
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS
AC021668/c
LOCUS
DEFINITION
AC021668
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS
1 (bases 1 to 43886)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
Anderson,S., Baldwin,J., Barna,N., Beckerly,R., Beda,F.,
Boguslavskiy,L., Boukhgalter,B., Brown,A., Burkett,G., Castle,A.,
Choepel,X., Colangelo,M., Collins,S., Collymore,A., Cooke,P.,
DeArellano,K., Dewar,K., Domino,M., Doyle,M., Fenestor,J.,
Ferreira,P., FitzHugh,W., Forrest,C., Gage,D., Galagan,J.,
Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L.,
Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J.,
Landers,T., Lechoczky,J., Levine,R., Liu,C., Liu,G., Locke,K.,
Macdonald,P., Marckay,N., McEwan,P., McGurk,A., McKernan,K.,
McPheeters,R., Meldrim,J., Meneus,L., Morrow,J., Naylor,J.,
Norman,C.H., O'Connor,T., O'Donnell,P., Oliver,T.M., Peterson,K.,
Pierre,N., Pisani,R., Pollara,V., Raymond,C., Riley,R., Rothman,D.,
Roy,A., Santos,R., Severy,P., Spencer,B., Stange-Thomann,N.,
Stojanovic,N., Subramanian,A., Talamas,J., Teefaye,S., Theodore,J.,
Tirrell,A., Vassiliev,H., Viel,R., Vo,A., Wu,X., Wyman,D., Ye,W.J.,
Zimmer,A. and Zody,M.
Direct Submission
Submitted (19-JAN-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Mar 29, 2001 this sequence version replaced gi:11612359.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L1276
Center clone name: 133_O_6
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* NOTE: This record contains 53 individual
* sequencing reads that have not been assembled into
* contigs. Runs of N are used to separate the reads
* and the order in which they appear is completely
* arbitrary. Low-pass sequence sampling is useful for
* identifying clones that may be gene-rich and allows
* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone
* will be sequenced to completion. In the event that
* the record is updated, the accession number will
* be preserved.
* 1
* 745: contig of 745 bp in length
* 846
* 845: gap of 100 bp
* 1576: contig of 731 bp in length
* 1677
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* 2410: contig of 734 bp in length
* 2511
* 2510: gap of 100 bp
* 3232: contig of 722 bp in length
* 3233
* 3232: gap of 100 bp
* 3333
* 4060: contig of 728 bp in length
* 4160: gap of 100 bp
* 4891: contig of 731 bp in length
* 4991: gap of 100 bp
* 5708: contig of 717 bp in length
* 5808: gap of 100 bp
* 5809: contig of 729 bp in length
* 6337: gap of 100 bp
* 6338
* 7344: contig of 707 bp in length
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* 8267: gap of 100 bp
* 8995: contig of 728 bp in length
* 9095: gap of 100 bp
* 9839: contig of 744 bp in length
* 9839: gap of 100 bp
* 9940
* 10653: contig of 714 bp in length
* 10753: gap of 100 bp
* 11497: contig of 744 bp in length
* 11498
* 11597: gap of 100 bp
* 12315: contig of 718 bp in length
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* 13136: contig of 721 bp in length
* 13236: gap of 100 bp
* 13367: contig of 731 bp in length
* 13968
* 14067: gap of 100 bp
* 14802: contig of 735 bp in length
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* 15642: contig of 740 bp in length
* 15742: gap of 100 bp
* 16477: contig of 735 bp in length
* 16577: gap of 100 bp
* 17301: contig of 724 bp in length
* 17401: gap of 100 bp
* 18129: contig of 728 bp in length
* 18229: gap of 100 bp
* 18958: contig of 729 bp in length
* 19058: gap of 100 bp
* 19787: contig of 729 bp in length
* 19788
* 20627: contig of 740 bp in length
* 20628
* 20727: gap of 100 bp
* 21449: contig of 722 bp in length
* 21549: gap of 100 bp
* 22275: contig of 726 bp in length
* 22375: gap of 100 bp
* 23103: contig of 728 bp in length
* 23203: gap of 100 bp
* 23939: contig of 736 bp in length
* 24039: gap of 100 bp
* 24775: contig of 736 bp in length
* 24875: gap of 100 bp
* 25602: contig of 727 bp in length
* 25702: gap of 100 bp
* 26418: contig of 716 bp in length
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* 27260: contig of 742 bp in length
* 27360: gap of 100 bp
* 28107: contig of 747 bp in length
* 28207: gap of 100 bp
* 28931: contig of 724 bp in length
* 29031: gap of 100 bp
* 29376: contig of 745 bp in length
* 29776: gap of 100 bp
* 30592: contig of 716 bp in length
* 30692: gap of 100 bp
* 31422: contig of 730 bp in length
* 31522: gap of 100 bp
* 32255: contig of 733 bp in length
* 32355: gap of 100 bp
* 33073: contig of 718 bp in length
* 33173: gap of 100 bp
* 33891: contig of 718 bp in length

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Oy      584  TAAATATAGAGCTTACTAGAAATTTGGCATCTAGAAATCTTGCTCCAAAGG 638
Db      134630 TAAATATAGAGCTTACTAGAAATTTGGCATCTAGAAATCTTGCTCCAAAGG 134576

RESULT 12 102(b)
AC016527/c
LOCUS      AC016527      208181 bp      DNA      linear      HTG 04-JUN-2000
DEFINITION Homo sapiens chromosome 15 clone RP11-460A24 map 15q21, LOW-PASS
SEQUENCE SAMPLING.
ACCESSION AC016527
VERSION   AC016527.2 GI:8247792
KEYWORDS HTG; HTGS_PHASE0.
SOURCE    Homo sapiens
ORGANISM  Homo sapiens
REFERENCE
AUTHORS   Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
          1 (bases 1 to 208181)
          Rowen,L., Madan,A., Qin,S., Abbasi,N., Baradarani,L., Birditt,B.,
          Bloom,S., Dors,M., Dickhoff,R., Fleetwood,P., Harrison,G.,
          Madan,A., Nesbitt,R., Shaffer,T. and Hood,L.
          Sequencing of human chromosome 15 D15S146-D15S117 region
          Unpublished
          2 (bases 1 to 208181)
          Rowen,L., Madan,A., Qin,S., Abbasi,N., Baradarani,L., Birditt,B.,
          Bloom,S., Dors,M., Dickhoff,R., Fleetwood,P., Harrison,G.,
          James,R., Kaur,A., Madan,A., Owen,M.P., Ratcliffe,A., Shaffer,T.
          and Hood,L.
          Direct Submission
          Submitted (02-DEC-1999) Multimegabase Sequencing Center, University
          of Washington, PO BOX 357730, Seattle, WA 98195, USA
          On Jun 4, 2000 this sequence version replaced gi:6503271.
          ----- Genome Center
          Center: Multimegabase Sequencing Center
          Web site: http://chroma.mbt.washington.edu/msg_www
          Contact: leerowen@systemsbiology.org
          ----- Summary Statistics
          Sequencing vector: pUC18; L08752
          Chemistry: Dye-terminator Big Dye; 90% of reads
          Chemistry: Dye-terminator Big Dye; 10% of reads
          Assembly program: Phrap; version 0.990399
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          * NOTE: This record contains 191 individual
          * sequencing reads that have not been assembled into
          * contigs. Runs of N are used to separate the reads
          * and the order in which they appear is completely
          * arbitrary. Low-pass sequence sampling is useful for
          * identifying clones that may be gene-rich and allows
          * overlap relationships among clones to be deduced.
          * However, it should not be assumed that this clone
          * will be sequenced to completion. In the event that
          * the record is updated, the accession number will
          * be preserved.
          *
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          * 1918 2890: contig of 973 bp in length
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          * 2991 3851: contig of 861 bp in length
          * 3852 3951: gap of unknown length
          * 3952 5018: contig of 1067 bp in length
          * 5019 5118: gap of unknown length
          * 5119 5991: contig of 873 bp in length
          * 5992 6091: gap of unknown length
          * 6092 7038: contig of 947 bp in length
          * 7039 7138: gap of unknown length
          * 7139 8021: contig of 883 bp in length
          * 8022 8121: gap of unknown length
          * 8122 9152: contig of 1031 bp in length
          * 9153 9252: gap of unknown length
          *
          * 10196: contig of 944 bp in length
          * 10296: gap of unknown length
          * 11211: contig of 915 bp in length
          * 11311: gap of unknown length
          * 11312 12220: contig of 909 bp in length
          * 12321 13554: contig of 1234 bp in length
          * 13555 14552: gap of unknown length
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          * 14853 15927: contig of 1175 bp in length
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          * 18021 18120: gap of unknown length
          * 18121 19061: contig of 941 bp in length
          * 19062 20206: contig of 1045 bp in length
          * 20207 21198: gap of unknown length
          * 21199 22311: contig of 1013 bp in length
          * 22312 23315: contig of 904 bp in length
          * 23316 24425: contig of 1010 bp in length
          * 24426 25448: gap of unknown length
          * 25449 26468: contig of 920 bp in length
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          * 37497 38516: contig of 920 bp in length
          * 38517 39598: contig of 982 bp in length
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          * 44093 45076: contig of 884 bp in length
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* 67454: contig of 858 bp in length
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* 69566: contig of 858 bp in length
* 69666: gap of unknown length
* 70597: contig of 931 bp in length
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* 71587: contig of 890 bp in length
* 71687: gap of unknown length
* 72595: contig of 908 bp in length
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* 73669: contig of 974 bp in length
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* 74685: contig of 916 bp in length
* 74785: gap of unknown length
* 75768: contig of 983 bp in length
* 75868: gap of unknown length
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* 76886: gap of unknown length
* 77743: contig of 857 bp in length
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* 79065: contig of 1222 bp in length
* 79165: gap of unknown length
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Matches 150; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

* 122 AGGTCAACTTCTCCATTTTATTTAGGACATTTCTTTATGCTCCCATAGACATGAGA 181
* 22658 AGGTCAACTTCTCCATTTTATTTAGGACATTTCTTTATGCTCCCATAGACATGAGA 22599
* 182 GCAGCAAGCAAGATTGGGAATGACACCATTTGTTACCTTGTATATAAAGAGCGG 241
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Db      22598 GCAGCAAGCAAGATTGGGAATGACACCATTTGTTACCTTGTATATAAAGAGCGG 22539
QY      242 CCATGTGGGAGACCTTTCTGTGACGGGTG 271
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RESULT 13
AY178583S2      147 bp DNA linear PRI 02-OCT-2003
LOCUS      Pan troglodytes EKN1 (EKN1) gene, exon 3.
DEFINITION      Pan troglodytes EKN1 (EKN1) gene, exon 3.
ACCESSION      AY178584
VERSION      AY178584.1 GI:27804579
KEYWORDS
SEGMENTS
SOURCE      2 of 9
ORGANISM      Pan troglodytes (chimpanzee)
Pan troglodytes
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
REFERENCE      1 (bases 1 to 147)
AUTHORS      Taipale,M., Kaminen,N., Nopola-Hemmi,J., Haltia,T., Myllyluoma,B.,
Lyytinen,H., Muller,K., Kaaranen,M., Lindsberg,P.J.,
Hannula-Jouppi,K. and Kere,J.
TITLE      A candidate gene for developmental dyslexia encodes a nuclear
tetrapeptide repeat domain protein dynamically regulated in
brain
JOURNAL      Proc. Natl. Acad. Sci. U.S.A. 100 (20), 11553-11558 (2003)
MEDLINE      22882828
PUBMED      12954984
REFERENCE      2 (bases 1 to 147)
AUTHORS      Kaminen,N. and Kere,J.
TITLE      EKN1 gene orthologs in chimpanzee, pygmy chimpanzee, gorilla and
orangutan
JOURNAL      Unpublished
REFERENCE      3 (bases 1 to 147)
AUTHORS      Kaminen,N. and Kere,J.
TITLE      Direct Submission
JOURNAL      Submitted (13-NOV-2002) Department of Medical Genetics, University
of Helsinki, PO Box 63, Haartmaninkatu 8, Helsinki FIN-00014,
Finland

FEATURES
source      1..147
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              /mol_type="genomic DNA"
              /db_xref="taxon:9598"
exon        1..147
              /gene="EKN1"
              /number=3

ORIGIN
Query Match      11.6%; Score 147; DB 9; Length 147;
Best Local Similarity 100.0%; Pred. No. 2.1e-63;
Matches 147; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      124 GTCAACTTCTCCATTTTATTTAGGACATTTCTTTATGCTCCCATAGACATGAGGC 183
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Db      1 GTCAACTTCTCCATTTTATTTAGGACATTTCTTTATGCTCCCATAGACATGAGGC 60
|||||
QY      184 AGCAAGCAAGATTGGGAATGACACCATTTGTTACCTTGTATATAAAGAGCGGC 243
|||||
Db      61 AGCAAGCAAGATTGGGAATGACACCATTTGTTACCTTGTATATAAAGAGCGGC 120
|||||
QY      244 ATGTGGGAGACCTTTCTGTGACGGGT 270
|||||
Db      121 ATGTGGGAGACCTTTCTGTGACGGGT 147
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RESULT 14
AY178583S5      147 bp DNA linear PRI 02-OCT-2003
LOCUS      Pan troglodytes EKN1 (EKN1) gene, exon 6.
DEFINITION      Pan troglodytes EKN1 (EKN1) gene, exon 6.
ACCESSION      AY178587
VERSION      AY178587.1 GI:27804582
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; CURRENT FILING DATE: 1999-02-12
; PRIOR APPLICATION NUMBER: US 60/074,725
; PRIOR FILING DATE: 1998-02-13
; PRIOR APPLICATION NUMBER: US 60/096,409
; PRIOR FILING DATE: 1998-08-13
; NUMBER OF SEQ ID NOS: 28208
; SEQ ID NO 12670
; LENGTH: 216
; TYPE: DNA
; ORGANISM: Candida albicans
US-09-248-796A-12670

Query Match 1.5%; Score 19; DB 4; Length 216;
Best Local Similarity 100.0%; Pred. No. 54;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 572 GAAGAAAAAAATAATAA 590
Db 63 GAAGAAAAAAATAATAA 81

RESULT 10
US-09-513-999C-11853/c
; Sequence 11853 Application US/09513999C
; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Duclert, A.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
; Patent No. 6783961
; FILE REFERENCE: 59.US2.REG
; CURRENT APPLICATION NUMBER: US/09/513,999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/122,487
; PRIOR FILING DATE: 1999-02-26
; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
; SEQ ID NO 11853
; LENGTH: 359
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 241
; OTHER INFORMATION: s-g or c
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 242
; OTHER INFORMATION: w-a or t
US-09-513-999C-11853

Query Match 1.5%; Score 19; DB 4; Length 359;
Best Local Similarity 100.0%; Pred. No. 54;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 570 AGGAAGAAAAAAATAAAA 588
Db 180 AGGAAGAAAAAAATAAAA 162

RESULT 11
US-08-343-443B-45/c
; Sequence 45, Application US/08343443B
; Patent No. 5968734
; GENERAL INFORMATION:
; APPLICANT: Aurias, Alain
; APPLICANT: Delattre, Olivier
; APPLICANT: Desmaze, Chantal
; APPLICANT: Melot, Thomas
; APPLICANT: Peter, Martine
; APPLICANT: Ploougastel, Beatrice
; APPLICANT: Thomas, Gilles

; APPLICANT: Zucman, Jessica
; TITLE OF INVENTION: NUCLEIC ACID CORRESPONDING TO A GENE OF
; TITLE OF INVENTION: CHROMOSOME 22 INVOLVED IN RECURRENT CHROMOSOMAL
; TITLE OF INVENTION: TRANSLATIONS ASSOCIATED WITH THE DEVELOPMENT OF CANCEROUS
; TITLE OF INVENTION: TUMORS, AND NUCLEIC ACIDS OF FUSION RESULTING FROM SAID
; NUMBER OF SEQUENCES: 129
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Weiser & Associates
; STREET: 230 South Fifteenth Street
; CITY: Philadelphia
; STATE: PA
; COUNTRY: USA
; ZIP: 19102
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: AEDIT 1.0 DOS text editor
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/343,443B
; FILING DATE: 18-NOV-1994
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/FR93/00494
; FILING DATE: 19-MAY-1993
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: FR 92/06123
; FILING DATE: 20-MAY-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Weiser, Gerard J.
; REGISTRATION NUMBER: 19,763
; REFERENCE/DOCKET NUMBER: 989.6121P
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 215-875-8383
; TELEFAX: 215-875-8394
; INFORMATION FOR SEQ ID NO: 45:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 425 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
US-08-343-443B-45

Query Match 1.5%; Score 19; DB 2; Length 425;
Best Local Similarity 100.0%; Pred. No. 54;
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 565 AAAGAAGGAAGAAAAAAA 583
Db 275 AAAGAAGGAAGAAAAAAA 257

RESULT 12
US-09-540-236-389
; Sequence 389, Application US/09540236
; Patent No. 6673910
; GENERAL INFORMATION:
; APPLICANT: Gary L. Breton et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO MORAXELLA CATAR
; TITLE OF INVENTION: FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 2709.2005-001
; CURRENT APPLICATION NUMBER: US/09/540,236
; CURRENT FILING DATE: 2000-04-04
; NUMBER OF SEQ ID NOS: 3840
; SEQ ID NO 389
; LENGTH: 507
; TYPE: DNA
; ORGANISM: M.catarrhalis
US-09-540-236-389

Query Match 1.5%; Score 19; DB 4; Length 507;
Best Local Similarity 100.0%; Pred. No. 54;

Matches	19;	Conservative	0;	Mismatches	0;	Indels	0;	Gaps	0;
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Db 383 TAAAGAAGACAAAAGAA 401

RESULT 13

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RESOLUTION 13
US-09-949-016-23803
; Sequence 23803, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 23803
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-23803

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Qy 290 AAAGAAATTAGAGAAAAATC 308
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pb 180 AAAGAAATTAGAGAAAAATC 198

PRECIPITATION

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RESULT 14
US-09-949-016-23804
; Sequence 23804, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 23804
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-23804

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QY 290 AAAGAATTAGAGAAAAATC 308
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b 422 AAAGAATTAGAGAAAAATC 440
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RESIST 15

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RESOLUTION IS
US-09-949-016-51133/c
; Sequence 51133, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 51133
; LENGTH: 601
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-51133

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Search completed: July 8, 2005, 12:07:45
Job time : 248 secs

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GenCore version 5.1.6

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OM nucleic - nucleic search, using sw model

Run on: July 8, 2005, 01:36:52 ; Search time 5659 Seconds

(without alignments)
10814.444 Million cell updates/sec

Title: US-10-681-199-1

Perfect score: 1263

Sequence: 1 atgctcttcaggttagcga.....gaacagaactaaatcttaa 1263

Scoring table:

IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 4708233 seqs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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GenEmbl.*

1: gb_ba.*

2: gb_hg.*

3: gb_in.*

4: gb_om.*

5: gb_ov.*

6: gb_pat.*

7: gb_ph.*

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9: gb_pr.*

10: gb_ro.*

11: gb_ats.*

12: gb_sy.*

13: gb_un.*

14: gb_vl.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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2	1047.4	82.9	1559	6 AX833451	AX833451 Sequence
3	1047.4	82.9	1559	9 AK095201	AK095201 Homo sapi
4	1047.2	82.9	1468	9 BC062564	BC062564 Homo sapi
5	846	67.0	1946	10 BC026462	BC026462 Mus muscu
6	485.2	38.4	2045	5 BC077575	BC077575 Xenopus l
7	432.6	34.3	489	6 AR413597	AR413597 Sequence
8	432.6	34.3	489	6 AX970431	AX970431 Sequence
9	432.6	34.3	489	6 BD109150	BD109150 EST and e
10	401.6	31.8	464	6 C0685075	C0685075 Sequence
11	390.2	30.9	1443	5 BC065881	BC065881 Danio rer
12	298	23.6	761	5 CR387522	CR387522 Gallus ga
13	275	21.8	1133	5 CR389904	CR389904 Gallus ga
14	241	19.1	114149	9 AC022083	AC022083 Homo sapi
15	239.4	19.0	43886	2 AC021668	AC021668 Homo sapi
16	239.4	19.0	151133	9 AC011355	AC011355 Homo sapi
17	234.6	18.6	208181	2 AC016527	AC016527 Homo sapi
18	227.8	18.0	231	9 AY178586	AY178586 Pan trogl
19	227.8	18.0	231	9 AY178595	AY178595 Pan panis

20	223	17.7	231	9	AY178601S4	AY178601 Gorilla g
21	219.8	17.4	231	9	AY178610S4	AY178613 Pongo pyg
22	178.8	14.2	313	6	AX915011	AX915011 Sequence
23	178.8	14.2	313	6	BD050544	BD050544 Sequence
24	162.8	12.9	164	6	AX918710	AX918710 Sequence
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26	151.4	12.0	153	9	AY178583S7	AY178589 Pan trogl
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28	151.4	12.0	153	9	AY178601S7	AY178607 Gorilla g
29	151.4	12.0	153	9	AY178610S7	AY178616 Pongo pyg
30	147	11.6	147	9	AY178583S5	AY178584 Pan trogl
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32	147	11.6	147	9	AY178592S2	AY178593 Pan panis
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34	147	11.6	147	9	AY178601S2	AY178602 Gorilla g
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36	147	11.6	147	9	AY178610S2	AY178611 Pongo pyg
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38	134.4	10.6	1187	6	CQ732459	CQ732459 Sequence
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ALIGNMENTS

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LOCUS AF337549 1993 bp mRNA linear PRI 15-JUN-2004
DEFINITION Homo sapiens EKN1 (EKN1) mRNA, complete cds.
ACCESSION AF337549
VERSION AF337549.1 GI:18478647
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 1993)
Taipale, M., Kaminen, N., Nopola-Hemmi, J., Haltia, T., Myllyluoma, B.,
Lyttinen, H., Muller, K., Kaaranen, M., Lindsberg, P. J.,
Hannula-Jouppi, K. and Kere, J.

TITLE A candidate gene for developmental dyslexia encodes a nuclear
tetrapeptide repeat domain protein dynamically regulated in
brain

JOURNAL Proc. Natl. Acad. Sci. U.S.A. 100 (20), 11553-11558 (2003)
MEDLINE 22882828
PUBMED 12954984
REFERENCE 2 (bases 1 to 1993)
AUTHORS Taipale, M. and Kere, J.
TITLE A gene disrupted by translocation breakpoint in chromosome 15q21
JOURNAL Unpublished
REFERENCE 3 (bases 1 to 1993)
AUTHORS Taipale, M. and Kere, J.

TITLE Direct Submission
JOURNAL Submitted (17-JAN-2001) Finnish Genome Center, University of
Helsinki, Tukholmankatu 2, Helsinki 00014, Finland
FEATURES
source
1. .1993
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/chromosome="15"
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1. .1993
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translocation breakpoint"
369. .1631

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Qy	181	AGCAGCAAGCAAGATTTGGGAATGACACCACTTGTCTTCACTTGTATATAAAAGAACGG	240
Db	223	AGCAGCAAGCAAGATTTGGGAATGACACCACTTGTCTTCACTTGTATATAAAAGAACGG	282
Qy	241	GCCATGTGGGAGACCCCTTTCTGTGACGGGTGTGACAAAGAGATGATGCAAGAAATTAGA	300
Db	283	GCCATGTGGGAGACCCCTTTCTGTGACGGGTGTGACAAAGAGATGATGCAAGAAATTAGA	342
Qy	301	GAATAATCTATTATCAAGCAACAGAGAGACCAAGAGCTTACAGAGCAAAAGCTCTCA	360
Db	343	GAATAATCTATTATCAAGCAACAGAGAGACCAAGAGCTTACAGAGCAAAAGCTCTCA	402
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Db	403	GCAAGCGGGAAGATCAAAAATACCACTTAAGTGTTCATGATGAAGATTGAAGAGAGAG	462
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Qy	601	ACTAGAAATTTGGCATCTAGAAATCTTCTCCAAAAGGGAGAAAATTCAGAAAAATATTT	660
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Qy	661	ACTGAGAGTTAAGGAAGACAGTATCTCTGCTCTCTGCTCTGTCGAGTATTAAAAATC	720
Db	703	ACTGAGAGTTAAGGAAGACAGTATCTCTGCTCTCTGCTCTGTCGAGTATTAAAAATC	762
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Qy	781	GAGTGGCTACACAAACAGCTGAGCAGCAGAGCAATGAATCTGACATAGCTGAACTT	840
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Db	1003	AATAAGATGCCACTATTGTTTGAACCGGCTGCTTCCACCTCAAAACTAAAAAATTA	1062
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Db	1063	CACAAGGCTATTGAAGATCTTCTTAAGGC	1091
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LOCUS			
DEFINITION Homo sapiens cDNA FLJ37882 fis, clone BRSTN2000536.			
ACCESSION AK095201			
VERSION AK095201.1 GI:21754405			
KEYWORDS oligo capping; fis (full insert sequence).			
SOURCE Homo sapiens (human)			

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

AUTHORS

1

Ota, T., Suzuki, Y., Nishikawa, T., Otsuki, T., Sugiyama, T., Irie, R., Wakamatsu, A., Hayaashi, K., Sato, H., Nagai, K., Kimura, K., Makita, H., Sekine, M., Obayashi, M., Nishi, T., Shibahara, T., Tanaka, T., Ishii, S., Yamamoto, J., Saito, K., Kawai, Y., Isono, Y., Nakamura, Y., Nagahari, K., Murakami, K., Yasuda, T., Iwayanagi, T., Wagatsuma, M., Shiratori, A., Sudo, H., Hosoiri, T., Kaku, Y., Kodaira, H., Kondo, H., Sugawara, M., Takahashi, M., Kanda, K., Kamihara, K., Yokoi, T., Furuya, T., Kikawa, E., Omura, Y., Abe, K., Kamiyama, K., Katsuya, N., Sato, K., Tanikawa, M., Yanazaki, M., Nishimura, K., Ishibashi, T., Yamashita, H., Murakawa, K., Fujimori, K., Tanai, H., Kimata, M., Watanabe, M., Hiraoka, S., Chiba, Y., Ishida, S., Ono, Y., Takiguchi, S., Watanabe, S., Yosida, M., Hotuta, T., Kusano, J., Kanehori, K., Takahashi-Fujii, A., Hara, H., Tanase, T., Nomura, Y., Toqiya, S., Komai, F., Hara, R., Takeuchi, K., Arita, M., Imose, N., Musahino, K., Yuuki, H., Oshima, A., Sasaki, N., Aotsuka, S., Yoshikawa, Y., Matsunawa, H., Ichihara, T., Shiohata, N., Sano, S., Moriya, S., Momiyama, H., Satoh, N., Takami, S., Terashima, Y., Suzuki, O., Nakagawa, S., Senoh, A., Mizoguchi, H., Goto, Y., Shimizu, F., Wakebe, H., Hishigaki, H., Watanabe, T., Sugiyama, A., Takemoto, M., Kawakami, B., Yamazaki, M., Watanabe, K., Kumagai, A., Itakura, S., Fukuzumi, Y., Fujimori, Y., Koniyama, M., Tashiro, H., Tanigami, A., Fujiwara, T., Ono, T., Yamada, K., Fujii, Y., Ozaki, K., Hirao, M., Ohmori, Y., Kawabata, A., Hikiji, T., Kobatake, N., Inagaki, H., Ikeda, Y., Okamoto, S., Okitani, R., Kawakami, T., Noguchi, S., Itoh, T., Shigetani, K., Senba, T., Matsumura, K., Nakajima, Y., Mizuno, T., Morinaga, M., Sasaki, M., Togaashi, T., Oyama, M., Hata, H., Watanabe, M., Komatsu, T., Mizuehima-Sugano, J., Sato, T., Shirai, Y., Takahashi, Y., Nakagawa, K., Okumura, K., Nagase, T., Nomura, N., Kikuchi, H., Masuho, Y., Yamashita, R., Nakai, K., Yada, T., Nakamura, Y., Ohara, O., Isogai, T. and Sugano, S. Complete sequencing and characterization of 21,243 full-length human cDNAs

Nat. Genet. 36 (1), 40-45 (2004)

14702039

2

JOURNAL

PUBMED

REFERENCE

AUTHORS

Kawakami, B., Sugiyama, A., Takemoto, M., Sugiyama, T., Irie, R., Otsuki, T., Sato, H., Wakamatsu, A., Ishii, S., Yamamoto, J., Isono, Y., Kawai-Hio, Y., Saito, K., Nishikawa, T., Kimura, K., Yamashita, H., Matsuo, K., Nakamura, Y., Sekine, M., Kikuchi, H., Kanda, K., Watanabe, M., Murakawa, K., Kanehori, K., Takahashi-Fujii, A., Oshima, A., Suzuki, Y., Sugano, S., Nagahara, K., Masuho, Y., Nagai, K. and Isogai, T.

NEDO human cDNA sequencing project

Unpublished

3 (bases 1 to 1559)

Isogai, T. and Yamamoto, J.

JOURNAL

AUTHORS

TITLE

JOURNAL

Submitted (04-JUL-2002) Takao Isogai, FLJ Project (HRI Team); 2-6-7 Kazusa-Kamatari, Kisarazu, Chiba 292-0812, Japan

(E-mail: genomics@hri.co.jp, Tel: 81-438-52-3975, Fax: 81-438-52-3986)

NEDO human cDNA sequencing project supported by Ministry of Economy, Trade and Industry of Japan; cDNA full insert sequencing: Research Association for Biotechnology (RAB); cDNA library construction: Helix Research Institute (HRI) (supported by Japan Key Technology Center etc.); 5'- & 3'-end one pass sequencing: RAB, HRI, and Biotechnology Center, National Institute of Technology and Evaluation; clone selection for full insert sequencing: HRI and RAB;

annotation: HRI and RAB.

FEATURES

source

1. 1559

Location/Qualifiers

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ORIGIN

Query Match 82.9%; Score 1047.4; DB 9; Length 1559;
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Matches 1048; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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Db	163	AAGTCAACTTCTCCATTTTATTTGAGGCATTTCTTTATGCTCCCATGACGATGAG	222
Qy	181	AGCAGCAAGCAAGATTGGGAATGACACCATTTGTCTTACCTTGTATATAAAAAAGCG	240
Db	223	AGCAGCAAGCAAGATTGGGAATGACACCATTTGTCTTACCTTGTATATAAAAAAGCG	282
Qy	241	GCCATGTGGGAGACCTTTCTGTGACGGGTGTTGACAAAGAGATGATGCAAGNATTAG	300
Db	283	GCCATGTGGGAGACCTTTCTGTGACGGGTGTTGACAAAGAGATGATGCAAGNATTAG	342
Qy	301	GAAATATCTATTTTACAAGCACAGAGAGACGAAAGAGCTACAGAGCAAAAGCTGCA	360
Db	343	GAAATATCTATTTTACAAGCACAGAGAGACGAAAGAGCTACAGAGCAAAAGCTGCA	402
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Qy	661	ACTAGAGATTAAAGGAAGACAGTATTCCTGCTCTCTGTTGTCAGTATTAATAATC	720
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Db	883	TGCGATTTAAAGACAGAGAAAAAGAACCCACAGATGGTTGAAGGATAAAGGAAACAATTG	942
Qy	901	TTTGCAACGAAAACTATTGGCAGCTATCAATGCATATATTTAGCCATAAGACTAAAT	960
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Db	1003	AATAAGATGCCACTATTGTATTGAAACGGGGCTGCTTGCACCTCAAAAACTTAA	1062
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RESULT 4	BC062564	1468 bp	mRNA	linear	PRI 28-NOV-2003
LOCUS	BC062564	1468 bp	mRNA	linear	PRI 28-NOV-2003
DEFINITION	Homo sapiens dyslexia susceptibility 1 candidate 1, mRNA (cDNA clone MGC:70618 IMAGE:5163101), complete cds.				
ACCESSION	BC062564				
VERSION	BC062564.1	GI:38565951			
KEYWORDS	MGC.				
SOURCE	Homo sapiens (human)				
ORGANISM	Homo sapiens				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.				
AUTHORS	1 (bases 1 to 1468)				
	Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G., Klausner, R.D., Collins, F.S., Wagner, L., Shennan, C.M., Schuler, G.D., Altschul, S.F., Zebberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K., Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsieh, F., Diatchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L., Stapleton, M., Soares, M.B., Bonaldo, M.F., Casavant, T.L., Scheetz, T.E., Brownstein, M.J., Usdin, T.B., Toshiyuki, S., Carninci, P., Prange, C., Raha, S., Loquellano, N.A., Peters, G.J., Abramson, R.D., Mullahy, S.J., Bosak, S.A., McEwan, P.J., McKernan, K.J., Malek, J.A., Gunaratne, P.H., Richards, S., Worley, K.C., Hale, S., Garcia, A.M., Gay, L.J., Hulyk, S.W., Villalón, D.K., Muzny, D.M., Sodergren, E.J., Lu, X., Gibbs, R.A., Fahey, J., Helton, E., Kettaman, M., Madan, A., Rodriguez, S., Sanchez, A., Whiting, M., Madan, A., Young, A.C., Shevchenko, Y., Bouffard, G.G., Blakesley, R.W., Touchman, J.W., Green, E.D., Dickson, M.C., Rodriguez, A.C., Grimwood, J., Schmutz, J., Myers, R.M., Butterfield, Y.S., Krzyzanski, M.I., Skalska, U., Snailus, D.E., Schnerch, A., Schein, J.E., Jones, S.J., and Marra, M.A., 2002. Generation and initial analysis of more than 15,000 full-length human and mouse cDNA sequences				
TITLE	Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)				
JOURNAL	22388257				
MEDLINE	12477932				
REFERENCE	2 (bases 1 to 1468)				
AUTHORS	Strausberg, R.				
TITLE	Direct Submission				
JOURNAL	Submitted (24-NOV-2003) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590, USA				
REMARK	NIH-MGC Project URL: http://mgc.nci.nih.gov				
COMMENT	Contact: MGC help desk Email: cgapsb@mail.nih.gov Tissue Procurement: Life Technologies, Inc. cDNA Library Preparation: Life Technologies, Inc. cDNA Library Arrayed by: The I.M.A.G.E. Consortium (ILNL) DNA Sequencing by: National Institutes of Health Intramural Sequencing Center (NISC), Gaithersburg, Maryland; Web site: http://www.nisc.nih.gov/ Contact: nisc_mgc@hgrl.nih.gov Akhtar, N., Ayale, K., Beckstrom-Sternberg, S.M., Benjamin, B., Blakesley, R.W., Bouffard, G.G., Breen, K., Brinkley, C., Brooks, S., Dietrich, N.L., Granite, S., Guan, X., Gupta, J., Haghighi, P., Hansen, N., Ho, S.-L., Karlins, E., Kwong, P., Laric, P., Legaspi, R.,				

Maduro, Q.L., Masiello, C., Maskeri, B., Mastrian, S.D., McCloskey, J.C., McDowell, J., Pearson, R., Scantripop, S., Thomas, P.J., Touchman, J.W., Tsourgeon, C., Vogt, J.L., Walker, M.A., Wetherby, K.D., Wiggins, L., Young, A., Zhang, L.-H. and Green, E.D.

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
Series: IRAK Plate: 135 Row: a Column: 3
This clone was selected for full length sequencing because it passed the following selection criteria: matched mRNA gi: 18677736.

FEATURES

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RESULT 5

BC026462 1946 bp mRNA linear ROD 29-JUN-2004
Mus musculus dyslexia susceptibility 1 candidate 1 homolog (human),
DEFINITION mRNA (cdna clone MGC:31465 IMAGE:4483015), complete cds.

ACCESSION

BC026462
GI:20071433

VERSION

BC026462.1

KEYWORDS

MGC.

SOURCE

Mus musculus

ORGANISM

Mus musculus (house mouse)

REFERENCE

1 (bases 1 to 1946)

AUTHORS

Strausberg, R.L., Feingold, B.A., Grouse, L.H., Derge, J.G., Klausner, R.D., Collins, F.S., Wagner, L., Shenmen, C.M., Schuler, G.D., Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K., Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Hsieh, P., Diatchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L., Stapleton, M., Soares, M.B., Bonaldo, M.F., Casavant, T.L., Scheetz, T.E., Brownstein, M.J., Usdin, T.B., Toshiyuki, S., Carninci, P., Prange, C., Raha, S.S., Loquellano, N.A., Peters, G.J., Abramson, R.D., Mullahy, S.J., Bosak, S.A., McEwan, P.J., McKernan, K.J., Malek, J.A., Gunaratne, P.H., Richards, S., Worley, K.C., Hale, S., Garcia, A.M., Gay, L.J., Hulyk, S.W., Villalon, D.K., Muzny, D.M., Sodergren, E.J., Lu, X., Gibbs, R.A., Fahey, J., Helton, E., Kettman, M., Madan, A., Rodriguez, S., Sanchez, A., Whiting, M., Madan, A., Young, A.C., Shevchenko, Y., Bouffard, G.G., Blakesley, R.W., Touchman, J.W., Green, E.D., Dickson, M.C., Rodriguez, A.C., Grimwood, J., Schmutz, J., Myers, R.M., Butterfield, Y.S., Krzywinska, M.I., Skalska, U., Smalusz, D.E., Schnerch, A., Schein, J.E., Jones, S.J. and Marra, M.A.

TITLE

Generation and initial analysis of more than 15,000 full-length

human and mouse cDNA sequences
 Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
 12477932
 2 (bases 1 to 1946)
 REFERENCE
 AUTHORS Strausberg, R.
 TITLE Direct Submission
 JOURNAL Submitted (02-APR-2002) National Institutes of Health, Mammalian
 Gene Collection (MGC), Cancer Genomics Office, National Cancer
 Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
 USA
 REMARK NIH-MGC Project URL: <http://mgc.nci.nih.gov>
 COMMENT Contact: MGC help desk
 Email: csapbe@mail.nih.gov
 Tissue Procurement: Gilbert Smith, Ph.D.
 cDNA Library Preparation: Life Technologies, Inc.
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Sequencing Group at the Stanford Human Genome
 Center, Stanford University School of Medicine, Stanford, CA 94305
 Web site: <http://www-shgc.stanford.edu>
 Contact: (Dickson, Mark) mcd@paxil.stanford.edu
 Dickson, M., Schmutz, J., Grimwood, J., Rodriguez, A., and Myers,
 R. M.
 Clone distribution: MGC clone distribution information can be found
 through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
 Series: IRAC Plate: 44 Row: b Column: 9
 This clone was selected for full length sequencing because it
 passed the following selection criteria: Hexamer frequency ORF
 analysis.
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DEFINITION	Xenopus laevis dyslexia susceptibility 1 candidate 1, mRNA (cDNA clone MGC:83582 IMAGE:5079663), complete cds.						

Schein, Asim Siddiqui, Rob Holt, Marco Marra.

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LINL at: <http://image.lnl.gov>

Series: IRAX Plate: 153 Row: a Column: 7

This clone was selected for full length sequencing because it passed the following selection criteria: Hexamer frequency ORF analysis. Similarity but not identity to protein.

FEATURES
SOURCE

gene

SDS

ORIGIN

Query Match	38.4%;	Score	485.2;	DB	5;	Length	2045;
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RESULT 7
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DEFINITION Sequence 1234 from patent US 6639063.
ACCESSION AR413597
VERSION AR413597.1 GI:40168707
KEYWORDS Unknown.
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 489)
AUTHORS Edwards,J.-B.D.M., Jobert,S. and Giordano,J.-Y.
TITLE EST's and encoded human proteins
JOURNAL Patent: US 6639063-A 1234 28-OCT-2003;
FEATURES Location/Qualifiers
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ORIGIN
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Qy 61 CTGCCCCCAAAGCGGTGCGTCAGAGACACGCGACGTGTTCTGCACGGAAACTATCTG 120
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Qy 361 GCAAAGCGGGAAGATCAAAATACGCACTAAGTGTCTATGATGAAGATTGAAGAAAGAGAG 420
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Qy 421 AGGAAAAAATAG 433
Db 477 AGGAAAAAATAG 489
RESULT 8
AX970431 LOCUS AX970431 489 bp DNA linear PAT 15-JAN-2004
DEFINITION Sequence 1234 from Patent EP1104808.
ACCESSION AX970431
VERSION AX970431.1 GI:40977781
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1
AUTHORS Dumas Milne Edwards,J.B., Jobert,S. and Giordano,J.Y.
TITLE ESTs and encoded human proteins
JOURNAL Patent: EP 1104808-A 1234 06-JUN-2001;
Genset (FR)
FEATURES Location/Qualifiers
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ORIGIN
Query Match 34.3%; Score 432.6; DB 6; Length 489;
Best Local Similarity 99.8%; Pred. No. 1.3e-74;
Matches 432; Conservative 1; Mismatches 0; Indels 0; Gaps 0;


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Qy 1 ATGCTCTTTCAGGTTAGCGATTACAGCTGGCAGCAGACGAAAGACTGCGGTCTTTCTGTCT 60
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Db 477 AGGAAAAAATAG 489

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RESULT 9
LOCUS BD109150 489 bp DNA linear PAT 18-SEP-2002
DEFINITION EST and encoded human protein.
ACCESSION BD109150
VERSION BD109150.1 GI:23203968
KEYWORDS JP 2002010789-A/1227.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
EDWARDS, J.B.D.M., Jobert, S. and Giordano, J.E.
EST and encoded human protein
Patent: JP 2002010789-A 1227 15-JAN-2002;
GENSET CORP
COMMENT OS Homo sapiens (human)
PN JP 2002010789-A/1227
PD 15-JAN-2002
PF 07-AUG-2000 JP 2000280989
PR 05-AUG-1999 US 60/147499
PI JEAN BAPTISTE DUMAS MILNE EDWARDS, SEVELIN JOBERT, JEAN EVE PI
GIORDANO
PC C12N15/09, C12N15/09, C07K14/47, C07K16/18, C12N1/15, C12N1/19, PC
C12N1/21,
PC C12N5/10, C12P21/02, C12P21/08, C12Q1/68, C12N15/00, C12N5/00, PC
C12N15/00
CC EST and encoded human protein
FH Key Location/Qualifiers
FT CDS 57..488.
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Query Match 34.3%; Score 432.6; DB 6; Length 489;
Best Local Similarity 99.8%; Pred. No. 1.3e-74;

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Qy 421 AGGAAAAAATAG 433
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RESULT 10
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DEFINITION Sequence 30001 from Patent WO02070737.
ACCESSION CQ685075
VERSION CQ685075.1 GI:42211607
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
LIEW, C.C., Marshall, W.E. and Zhang, H.
Compositions and methods relating to osteoarthritis
Patent: WO 02070737-A 30001 12-SEP-2002;
Chondrogene Inc. (CA)
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Matches 404; Conservative 0; Mismatches 7; Indels 0; Gaps 0;
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Qy 121 AAGGTCAACTTTCCTCCATTTTATTTGAGGCAATTTCTTTATGTCTCCCATAGACGATGAG 180
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181 AGCAGCAAGCAAGATTGGGAATGACACCATCTCTTACCTTGTTATATAAAGAGCG 240
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RESULT 11
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 ACCESSION BC065881
 VERSION BC065881.1 GI:41389019
 KEYWORDS MGC.
 SOURCE Danio rerio (zebrafish)
 ORGANISM Danio rerio

REFERENCE
 AUTHORS Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G., Klausner, R.D., Collins, F.S., Wagner, L., Shenmen, C.M., Schuler, G.D., Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K., Hopkins, R.F., Jordan, H., Moore, T., Max, S.I., Wang, J., Haieh, P., Diatchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L., Stapleton, M., Soares, M.B., Bonaldo, M.P., Casavant, T.L., Scheetz, T.E., Brownstein, M.J., Ustin, T.B., Toshiyuki, S., Carninci, P., Prange, C., Raha, S.S., Loquellano, N.A., Peters, G.J., Abramson, R.D., Mullahy, S.J., Bosak, S.A., McEwan, P.J., McKernan, K.J., Malek, J.A., Gunaratne, P.H., Richards, S., Worley, K.C., Hale, S., Garcia, A.M., Gay, L.J., Hulyk, S.W., Villalón, D.K., Muzny, D.M., Sodergren, E.J., Lu, X., Gibbs, R.A., Fahey, J., Helton, E., Kettman, M., Madan, A., Rodriguez, S., Sanchez, A., Whitting, M., Madan, A., Young, A.C., Shevchenko, Y., Bouffard, G.G., Blakesley, R.W., Touchman, J.W., Green, E.D., Dickson, M.C., Rodriguez, A.C., Grimwood, J., Schmutz, J., Myers, R.M., Butterfield, Y.S., Krzywinski, M.I., Skalka, U., Smailus, D.E., Schnerch, A., Schein, J.E., Jones, S.J. and Marra, M.A.
 Generation and initial analysis of more than 15,000 full-length human and mouse cDNA sequences
 Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
 12477932
 2 (bases 1 to 1443)

TITLE Strausberg, R.
 JOURNAL Direct Submission
 PUBMED Submitted (26-JAN-2004) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590, USA
 REFERENCE NIH-MGC Project URL: <http://mgc.nci.nih.gov>
 COMMENT Contact: MGC help desk
 Email: cgapbs-remail.nih.gov
 Tissue Procurement: Len Zon, Harvard
 cDNA Library Preparation: Open Biosystems
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Genome Sequence Centre, BC Cancer Agency, Vancouver, BC, Canada
 info@bcgsc.bc.ca
 Steve Jones, Sarah Barber, Mabel Brown-John, Yaron Butterfield, Andy Chan, Steve S. Chand, William Chow, Allison Cloutier, Ruth Featherstone, Malachi Griffith, Obi Griffith, Ran Guin, Nancy Liao,

Kim MacDonald, Amara Masson, Mike R. Mayo, Josh Moran, Ryan Morin, Teika Olson, Diana Palmquist, Anca Petrescu, Anna Liisa Prabbu, Parvaneh Saeedi, JR Santos, Angélique Schnerch, Ursula Skalka, Duane Smailus, Jeff Stott, Miranda Tsai, George Yang, Jacque Schein, Asim Siddiqui, Rob Holt, Marco Marra.

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
 Series: IRAK Plate: 147 Row: p Column: 24
 This clone was selected for full length sequencing because it passed the following selection criteria: Hexamer frequency ORF analysis, Similarity but not identity to protein.

FEATURES

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DEFINITION Gallus gallus finished cdna, clone CHEST505n19.
ACCESSION CR387522
VERSION CR387522.1 GI:46240281
KEYWORDS
SOURCE Gallus gallus (chicken)
ORGANISM Gallus gallus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
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Archosauria; Aves; Neognathae; Galliformes; Phasianidae;
Phasianinae; Gallus.
1 (bases 1 to 761)
Boardman,P.E., Bonfield,J.K., Brown,W.R.A., Carder,C., Chalk,S.E.,
Croning,M.D.R., Davies,R.M., Francis,M.D., Grafham,D.V.,
Hubbard,S.J., Humphray,S.J., Hunt,P.J., Maddison,M., McLaren,S.R.,
Niblett,D., Overton,I.M., Rogers,J., Scott,C.E., Taylor,R.G.,
Tickle,C. and Wilson,S.A.
Direct Submission
Submitted (05-APR-2004) Sanger Institute, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: chicken@bms.umist.ac.uk
BBSRC/Dundee/Nottingham/Sanger/Sheffield/UMIST Gallus gallus cdna
sequencing project.
This sequence is from the
BBSRC/Dundee/Nottingham/Sanger/Sheffield/UMIST cdna collection,
from a library constructed by Elizabeth Bosch. cdna was prepared
from RNA extracted from hearts, normalised, and poly A-tailed.
ECORI-NotI cut cdna was then ligated into the vector. Vector:
pBluescript II KS(+); Site_1: EcoRI; Site_2: NotI Host: Escherichia
coli DH10B.
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Best Local Similarity 76.8%; Pred. No. 2.8e-48;
Matches 364; Conservative 0; Mismatches 110; Indels 0; Gaps 0;
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CR389904
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DEFINITION Gallus gallus finished cdna, clone CHEST271n13.
ACCESSION CR389904
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VERSION CR389904.1 GI:46428549
KEYWORDS
SOURCE
ORGANISM Gallus gallus (chicken)
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REFERENCE 1 (bases 1 to 1133)
AUTHORS Boardman,P.E., Bonfield,J.K., Brown,W.R.A., Carder,C., Chalk,S.E.,
          Croning,M.D.R., Davies,R.M., Francis,M.D., Grafham,D.V.,
          Hubbard,S.J., Humphray,S.J., Hunt,P.J., Maddison,M., McLaren,S.R.,
          Niblett,D., Overton,I.M., Rogers,J., Scott,C.E., Taylor,R.G.,
          Tickle,C. and Wilson,S.A.
TITLE Direct Submission
JOURNAL Submitted (19-APR-2004) Sanger Institute, Hinxton, Cambridgeshire,
COMMENT CB10 ISA, UK. E-mail enquiries: chickest@ms.umbist.ac.uk
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          This sequence is from the
          BBSRC/Dundee/Nottingham/Sanger/Sheffield/UMIST cDNA collection,
          from a library constructed by Elizabeth Bosch. cDNA was prepared
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Best Local Similarity 66.4%; Pred. No. 8.8e-44;
Matches 395; Conservative 0; Mismatches 200; Indels 0; Gaps 0;
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Db 73 ATGCGCTGTGGCTGCGGAGTACAGCTGGCGGAGCGCGCGGTACCTCTCG 132
Qy 61 CTGCCCTTCAAGCGGTGTGCGTCAGAGACACGAGCGTGTCTGCAGGAAACATCTGTG 120
Db 133 CTGCCCGTGCAGCGCTGAGGCTCACCGCCGCCAACATCTTCTGCACGCGGTACCTG 192
Qy 121 AAGTCACCTTCTCCATTTTATTTGAGGCATTTCTTTATGCTCCCATAGACGATGAG 180
Db 193 AAGGTAGCGTCCCTCCCTTTTATTCGAGCTGTCTGTAGTCTCTTATTTGATGAGACA 252
Qy 181 AGCAGCAAGCAAGATTGGGAATGACACCAATGCTTCTTACCTTGTATATAAAGAGCG 240
Db 253 AATAGCAGCAAGATCGCAATGGAGTGTCTTCTTCTTCTGATATAAAGAGAGCG 312
Qy 241 GCCATGTGGAGACCTTTCTGTGAGCGGTGTGCAAGAGATGATGCAAGAAATTAGA 300
Db 313 GCCATGTGGAGTCCCTGGCTGTGCAAAATGTTAACAGAGGAAGCACTGCAACGCTAAGA 372
Qy 301 GAAAAATCTATTTACAGACACAGAGAGCAAAAGAGCTACAGAGCAAAAGCTGCA 360
Db 373 GAGAAATGCTGTCTGAAAGACACAGAAAGCAAAAGAGGAGAGAGCAAAAGTT 432
Qy 361 GCAAAGCGGGAAGATCAAAAATAGCGCACTAAGTGTGTCATGATGAAGATTGAAGAGAGAG 420
Db 433 ACAAAACAGGAACATAGAAGTATGCTTTTGAGGCTACTAATGAAGCTAGAGAGAGAGAG 492
Qy 421 AGGAAAAAATAGAGATATGAAGAAATGAACGGATATAAGCCACTAAAGCTTTGGAA 480
Db 493 AGAAAAAGAAATGAAGATCTGAAAGAAACAGAGAGACAGCAAGTGCCTAAGGAGTTGGAG 552
Qy 481 GCCTGGAAAGAAATATCAAGAAAGAAAGCTGAGGAGCAAAAAAATTCAGAGAGAGAGAGAA 540
Db 552
Qy 543 CTATGAGAGAGTCAAGAAACGAGTGTGAGAAACAAAGAGGCTACAAAAGAGGAGAA 612
Qy 541 TTATGTCAAAAGAAAGCAATTAAGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 595
Db 613 CTACATGAAGAGTAGAGCACTAAAGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAG 667
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LOCUS Homo sapiens chromosome 15 clone CTD-2137J4 map 15q21.3, complete
DEFINITION sequence.
AC022083 GI:14029048
VERSION AC022083.6
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Eukaryota; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 114149)
AUTHORS Rowen,L., Madan,A., Qin,S., Baradarani,L., Birditt,B., Bloom,S.,
          Burke,J., Dors,M., Fleetwood,P., Kaur,A., Madan,A., Nesbitt,R.,
          Pate,D. and Hood,L.
TITLE Sequencing of human chromosome 15 D15S146-D15S117 region
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 114149)
AUTHORS Rowen,L., Madan,A., Qin,S., Abbasi,N., Baradarani,L., Birditt,B.,
          Bloom,S., Dors,M., Dickhoff,R., Fleetwood,P., Harrison,G., James,R.,
          Kaur,A., Madan,A., Owen,M.P., Ratcliffe,A., Shaffer,T. and Hood,L.
TITLE Direct Submission
JOURNAL Submitted (26-JAN-2000) Multimegabase Sequencing Center, University
          of Washington, PO BOX 357730, Seattle, WA 98195, USA
REFERENCE 3 (bases 1 to 114149)
AUTHORS Rowen,L., Madan,A., Qin,S., Baradarani,L., Birditt,B., Bloom,S.,
          Burke,J., Dors,M., Fleetwood,P., Kaur,A., Madan,A., Nesbitt,R.,
          Pate,D. and Hood,L.
TITLE Direct Submission
JOURNAL Submitted (12-MAY-2001) Multimegabase Sequencing Center, Institute
          for Systems Biology, 4225 Roosevelt Way NE, Suite 200, Seattle, WA
          98105, USA
COMMENT On May 12, 2001 this sequence version replaced gi:13621222.
----- Genome Center
Center: Multimegabase Sequencing Center
Center code: UWMSC
Web site: http://chroma.mbt.washington.edu/msg_www
Contact: leerowensystemsbiology.org
----- Summary Statistics
Sequencing vector: pUC18; L08752
Chemistry: Dye-terminator Big Dye; 90% of reads
Chemistry: Dye-primer Big Dye; 10% of reads
Assembly program: Phrap; version 0.990399
Note: Data from overlapping clones AC013355 [Drafting center:
UWMSC], AC012674 [Drafting center: BCM] and AC012378 [Drafting
center: UWMSC] were added for finishing.
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                     /clone="CTD-2137J4"
                     /clone_lib="Cal Tech human BAC library D"
                     /note="This clone overlaps RP11-178D12 AC013355,
                     RP11-458H3 AC012674 and RP11-420M1 AC012378. Data from
                     overlapping BACs were added and the consensus sequence
                     determined from CTD-2137J4 to the extent possible."
     misc_feature      1..47896
                     /note="overlap with RP11-178D12, AC013355"
     misc_feature      18530..18640
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95860..114149
/note="overlap with RP11-458H3, AC012674"
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/note="overlap with RP11-420M1, AC012378"
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ORIGIN

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31582 TGTATAGATTGAAGAAGAGAGAGGAAAAAATAGAAGATATGAAGAATAATGAACGGA 31523
Qy 458 TAAAGCCACTTAAGCATTGGAGCTGGAAGAATATCAAGAAGAAAGCTGAGGAGCAAA 517
Db |||
31522 TAAAGCCACTTAAGCATTGGAGCTGGAAGAATATCAAGAAGAAAGCTGAGGAGCAAA 31463
Qy 518 AAAAATTCAGAGAGAGAGAAATATGTCAAAAAGAAAGCAAAATTAAGAAGCAAGAA 577
Db |||
31462 AAAAATTCAGAGAGAGAGAAATATGTCAAAAAGAAAGCAAAATTAAGAAGCAAGAA 31403
Qy 578 AAAAATTAATATGAAGTCTTACTAGAAATTTGGCATCTAGAATCTTGCTCCAAAAG 637
Db |||
31402 AAAAATTAATATGAAGTCTTACTAGAAATTTGGCATCTAGAATCTTGCTCCAAAAG 31343
Qy 638 GGAGAAATTCAGAAATATATTTACTGAGAAGT 670
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RESULT 15
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DEFINITION Homo sapiens chromosome 15 clone RP11-13306 map 15, LOW-PASS
SEQUENCE SAMPLING.
AC021668
VERSION AC021668.4 GI:13488020
KEYWORDS HTG; HTGS_PHASE0.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 43886)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Abrahams,H., Allen,N.,
Anderson,S., Baldwin,J., Barna,N., Becker,R., Bieda,F.,
Boguslavsky,L., Boukhgalter,B., Brown,A., Burkett,G., Castle,A.,
Choepel,Y., Colangelo,M., Collins,S., Collymore,A., Cooke,P.,
Dearellano,K., Dewar,K., Domino,M., Doyle,M., Fenebor,J.,
Ferreira,P., FitzHugh,W., Forrest,C., Gage,D., Galagan,J.,
Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L.,
Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J.,
Landers,T., Lechoczy,J., Levine,R., Liu,C., Liu,G., Locke,K.,
Macdonald,P., Marquis,N., McEwan,P., McGurk,A., McKernan,K.,
McPheeters,R., Meldrim,J., Meneus,L., Morrow,J., Naylor,J.,
Norman,C.H., O'Connor,T., O'Donnell,P., Olivar,T.M., Peterson,K.,
Pierre,N., Pisan,C., Pollara,V., Raymond,C., Riley,R., Rothman,D.,
Roy,A., Santos,R., Severy,P., Spencer,B., Stange-Thomann,N.,
Stojanovic,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
Tirrell,A., Vassiliev,H., Viel,R., Vo,A., Wu,X., Wyman,D., Ye,W.J.,
Zimmer,A. and Zody,M.
Direct Submission
Submitted (19-JAN-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA

```

COMMENT

On Mar 29, 2001 this sequence version replaced gi:11612359.
 All repeats were identified using RepeatMasker:
 Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>
 ----- Genome Center
 Center: Whitehead Institute/ MIT Center for Genome Research
 Center code: WIBR
 Web site: <http://www-seq.wi.mit.edu>
 Contact: sequence_submissions@genome.wi.mit.edu
 ----- Project Information
 Center project name: L1276
 Center clone name: 133_O_6

 * NOTE: This record contains 53 individual
 * sequencing reads that have not been assembled into
 * contigs. Runs of N are used to separate the reads
 * and the order in which they appear is completely
 * arbitrary. Low-pass sequence sampling is useful for
 * identifying clones that may be gene-rich and allows
 * overlap relationships among clones to be deduced.
 * However, it should not be assumed that this clone
 * will be sequenced to completion. In the event that
 * the record is updated, the accession number will
 * be preserved.
 * 1 745: contig of 745 bp in length
 * 746 845: gap of 100 bp
 * 846 1576: contig of 731 bp in length
 * 1577 1676: gap of 100 bp
 * 1677 2410: contig of 734 bp in length
 * 2411 2510: gap of 100 bp
 * 2511 3232: contig of 722 bp in length
 * 3233 3332: gap of 100 bp
 * 3333 4060: contig of 728 bp in length
 * 4061 4160: gap of 100 bp
 * 4161 4891: contig of 731 bp in length
 * 4892 4991: gap of 100 bp
 * 4992 5708: contig of 717 bp in length
 * 5709 5808: gap of 100 bp
 * 5809 6537: contig of 729 bp in length
 * 6538 6637: gap of 100 bp
 * 6638 7344: contig of 707 bp in length
 * 7345 7444: gap of 100 bp
 * 7445 8167: contig of 723 bp in length
 * 8168 8267: gap of 100 bp
 * 8268 8995: contig of 728 bp in length
 * 8996 9095: gap of 100 bp
 * 9096 9839: contig of 744 bp in length
 * 9840 9939: gap of 100 bp
 * 9940 10653: contig of 714 bp in length
 * 10654 10753: gap of 100 bp
 * 10754 11497: contig of 744 bp in length
 * 11498 11597: gap of 100 bp
 * 11598 12315: contig of 718 bp in length
 * 12316 12415: gap of 100 bp
 * 12416 13136: contig of 721 bp in length
 * 13137 13236: gap of 100 bp
 * 13237 13967: contig of 731 bp in length
 * 13968 14067: gap of 100 bp
 * 14068 14802: contig of 735 bp in length
 * 14803 14902: gap of 100 bp
 * 14903 15642: contig of 740 bp in length
 * 15643 15742: gap of 100 bp
 * 15743 16477: contig of 735 bp in length
 * 16478 16577: gap of 100 bp
 * 16578 17301: contig of 724 bp in length
 * 17302 17401: gap of 100 bp
 * 17402 18129: contig of 728 bp in length
 * 18130 18229: gap of 100 bp
 * 18230 18958: contig of 729 bp in length
 * 18959 19058: gap of 100 bp
 * 19059 19787: contig of 729 bp in length
 * 19788 19887: gap of 100 bp
 * 19888 20627: contig of 740 bp in length

GenCore version 5.1.6
Copyright (c) 1993 - 2005 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 8, 2005, 01:36:17 ; Search time 755 Seconds
(without alignments)
9902.826 Million cell updates/sec

Title: US-10-681-199-1
Perfect score: 1263
Sequence: 1 atgctcttcaggtagcgca.....gaacagaactaaatcttaa 1263

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N_Geneseq_16Dec04:*
1: Geneseqn1980s:*
2: Geneseqn1990s:*
3: Geneseqn2000s:*
4: Geneseqn2001as:*
5: Geneseqn2001bs:*
6: Geneseqn2002as:*
7: Geneseqn2002bs:*
8: Geneseqn2003as:*
9: Geneseqn2003bs:*
10: Geneseqn2003cs:*
11: Geneseqn2003ds:*
12: Geneseqn2004as:*
13: Geneseqn2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
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2	1263	100.0	1263	9	ADB16965 cDNA sequ
3	1253.4	99.2	1263	9	ADB16939 Pygmy chi
4	1251.8	99.1	1263	9	ADB16933 Chimpanze
5	1248.6	98.9	1263	9	ADB16935 Gorilla D
6	1243.8	98.5	1263	9	ADB16937 Orangutan
7	1049	83.1	1641	10	ADC30210 Human nov
8	1047.4	82.9	1559	11	ADM01890 Human CDN
9	840.8	66.6	1697	9	ADB16924 cDNA sequ
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13	245.2	19.4	1383	5	AAS70018 DNA encod
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15	178.8	14.2	313	3	AAC26799 Human sec
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28	92.8	7.3	627	13	ACN54555	ACN54555 Cotton an
29	92.8	7.3	6668	6	ABL33697	Ab133697 Human imm
30	92.8	7.3	17934	6	ABL33719	Ab133719 Human imm
31	92.2	7.3	556	6	ABQ36997	Abq36997 Oligonucl
32	92.2	7.3	556	6	ABQ36996	Abq36996 Oligonucl
33	92.2	7.3	16633	6	ABN79984	Abn79984 Human che
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35	91.4	7.2	1000	12	ADQ62832	Adq62832 Homopoly-
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38	89.8	7.1	2226	12	ADI43251	Adi43251 Plant tra
39	89.8	7.1	2226	12	ADO02941	Ado02941 Soybean o
40	89.6	7.1	34769	4	AAS46774	Aas46774 Tumour su
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ALIGNMENTS

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ID ADB16964 standard; cDNA; 1263 BP.
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AC ADB16964;
XX
DT 20-NOV-2003 (first entry)
XX
DE Human DYXC1 cDNA with single nucleotide polymorphisms.
XX
KW Gene; ss; human; DYXC1; dyslexia; neurological disorder;
KW Chromosome 15q21; reading disability; phonological processing;
KW rapid naming; verbal short-term memory; single nucleotide polymorphism;
KW SNP.
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OS Homo sapiens.
XX
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WO2003068814-A1.
XX
PD 21-AUG-2003.
XX
PF 12-FEB-2003; 2003WO-FI000110.
XX
PR 12-FEB-2002; 2002US-0355782P.
XX
PA (LICN) LICENTIA LTD.

XX
PI Kere J, Taipale M, Nopola-Hemmi J, Kaminen N;
XX WPI; 2003-646482/61.
DR P-PSDB; ADB16923.
XX
XX New isolated, purified DYX1C1 nucleic acid for studying brain processes,
PT e.g. reading, phonological processing, rapid naming or verbal short-term
PT memory, or for diagnosing dyslexia or assessing the predisposition to
PT dyslexia.
XX
XX Claim 1; Page 47; 135pp; English.
XX
XX This invention relates to a novel isolated human gene DYX1C1 that is
CC functionally related to dyslexia, more particularly it describes single
CC nucleotide polymorphisms thought to predispose an individual in to
CC developing dyslexia. This is a neurological disorder with a genetic basis
CC (DYX1C1 has been isolated to chromosome 15q21), which manifests itself as
CC a specific reading disability. Specifically, DYX1C1 is can be useful in
CC study of brain processes such as reading, phonological processing, rapid
CC naming and verbal short-term memory. Accordingly, the present invention
CC describes methods and materials for analysing allelic variations in the
CC DYX1C1 gene, and also provides DYX1C1 as an antigen for the production of
CC antibodies used in the diagnosis of dyslexia. This polynucleotide
CC represents the coding sequence of human DYX1C1 cDNA of the invention.
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Best Local Similarity 100.0%; Pred. No. 1.3e-263;
Matches 1263; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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DB 1021 CACAAGCTATTTGAAGATTTCTTAAGGCACTGGAATTTATGATGCCACCTGTTACAGAC 1080
QY 1081 AATGCTTAATGCAAGATGAGGACACATGTACGACGTCGGAACAGCATTTCTGCACTAGAA 1140
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QY 1141 TTGTATGTAGAACGCTTACAGGATTTATGAACGCGCACTTAAGATTTGATCCATCCAAACAA 1200
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QY 1261 TAA 1263
DB 1261 TAA 1263
RESULT 2
ADBI6965
ID ADBI6965 standard; cDNA; 1993 BP.
XX
AC ADBI6965;
XX AC
XX AC
DT 20-NOV-2003 (first entry)
XX
DE cDNA sequence of the human DYX1C1 mRNA.
XX
XX gene; ss; human; DYX1C1; dyslexia; neurological disorder;
KW chromosome 15q21; reading disability; phonological processing;
KW rapid naming; verbal short-term memory; SNP;
KW single nucleotide polymorphism.
XX
OS Homo sapiens.
XX
XX Key Location/Qualifiers
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FT /standard_name= "Single nucleotide polymorphism"
FT 369..1631
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XX

PN WO2003068814-A1.
 XX 21-AUG-2003.
 XX 12-FEB-2003; 2003WO-FI000110.
 XX 12-FEB-2002; 2002US-0355782P.
 XX (LINC) LICENTIA LTD.
 XX Kere J, Taipale M, Nopola-Hemmi J, Kaminen N;
 XX WPI; 2003-646482/61.
 XX P-PsDB; ADB16933.
 DR New isolated, purified DYXC1 nucleic acid for studying brain processes,
 PT e.g. reading, phonological processing, rapid naming or verbal short-term
 PT memory, or for diagnosing dyslexia or assessing the predisposition to
 PT dyslexia.
 XX
 PS Claim 10; Page 48-50; 135pp; English.
 XX
 CC This invention relates to a novel isolated human gene DYXC1 that is
 CC functionally related to dyslexia, more particularly it describes single
 CC nucleotide polymorphisms thought to predispose an individual in to
 CC developing dyslexia. This is a neurological disorder with a genetic basis
 CC (DYXC1 has been isolated to chromosome 15q21), which manifests itself as
 CC a specific reading disability. Specifically, DYXC1 is can be useful in
 CC study of brain processes such as reading, phonological processing, rapid
 CC naming and verbal short-term memory. Accordingly, the present invention
 CC describes methods and materials for analysing allelic variations in the
 CC DYXC1 gene, and also provides DYXC1 as an antigen for the production of
 CC antibodies used in the diagnosis of dyslexia. This polynucleotide is the
 CC cDNA sequence of the wild type human DYXC1 mRNA of the invention.
 XX
 SQ Sequence 1993 BP; 693 A; 368 C; 430 G; 502 T; 0 U; 0 Other;
 Query Match 100.0%; Score 1263; DB 9; Length 1993;
 Best Local Similarity 100.0%; Pred. No. 1.4e-263;
 Matches 1263; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 1 ATGCTCTTCTAGGTTAGCGATTACAGCTGGCGAGGAGAGAGAGCTGGCGTCTTCTGTCT 60
 369 ATGCTCTTCTAGGTTAGCGATTACAGCTGGCGAGGAGAGAGAGCTGGCGTCTTCTGTCT 428
 61 CTGCCCCCTCAAGGGCGTGTGGCTCAGAGACACGAGCGTGTCTGACCGGAAACTATCTG 120
 429 CTGCCCCCTCAAGGGCGTGTGGCTCAGAGACACGAGCGTGTCTGACCGGAAACTATCTG 488
 121 AAGGTCAACTTTCCTCCATTTTATTTAGGCGATTTCTTTATGCTCCCATAGACGATGAG 180
 489 AAGGTCAACTTTCCTCCATTTTATTTAGGCGATTTCTTTATGCTCCCATAGACGATGAG 548
 181 AGCAGCAAGCAAGATTGGGATGACACCATTTGCTTCCCTGATATAAAGAAAGCGG 240
 549 AGCAGCAAGCAAGATTGGGATGACACCATTTGCTTCCCTGATATAAAGAAAGCGG 608
 241 GCCATGTGGGAGACCTTTCTGTGACGGGTGTGACAAAGAGATGATCAAGAAATTAGA 300
 609 GCCATGTGGGAGACCTTTCTGTGACGGGTGTGACAAAGAGATGATCAAGAAATTAGA 668
 301 GAAAAATCTATTTTCAAGCAACAGAGAGAGCAAAAGAGCTTACAGAGCAAAAGCTGCA 360
 669 GAAAAATCTATTTTACAGCAACAGAGAGAGCAAAAGAGCTTACAGAGCAAAAGCTGCA 728
 361 GCAAGCGGAGAGATCAAAATATCCGACTAAGTGTGATGATGAAGATTGAAGAAAGAG 420
 729 GCAAGCGGAGAGATCAAAATATCCGACTAAGTGTGATGATGAAGATTGAAGAAAGAG 788
 421 AGGAAAAAATAGAGATATGAAGAAATTAACCGATTAAGCAACCTAAAGCATTTGAA 480
 789 AGGAAAAAATAGAGATATGAAGAAATTAACCGATTAAGCAACCTAAAGCATTTGAA 848

QY 481 GCCTGGAAAGAAATATCAAGAAAGCTGAGAGCAAAAGAAATTTACAGAGAGAGAGAAA 540
 DB 849 GCCTGGAAAGAAATATCAAGAAAGCTGAGAGCAAAAGAAATTTACAGAGAGAGAGAAA 908
 QY 541 TTATGTCAAAAGAAAGCAAAATTAAGAAAGCAAAAGAAATTTAAATATAAAGAGTCTT 600
 DB 909 TTATGTCAAAAGAAAGCAAAATTAAGAAAGCAAAAGAAATTTAAATATAAAGAGTCTT 968
 QY 601 ACTAGAAATTTGGCATCTAGAAATCTTCTCCAAAGGGAGAAATTTAGAAATATATTTT 660
 DB 969 ACTAGAAATTTGGCATCTAGAAATCTTCTCCAAAGGGAGAAATTTAGAAATATATTTT 1028
 QY 661 ACTGAGAGTTTAAAGGAGAGACAGTATCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 720
 DB 1029 ACTGAGAGTTTAAAGGAGAGACAGTATCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 1088
 QY 721 AACTTTACCCCTCCAGATTTCCCAACAGCTCTTCTGTGAATCAACAAGTACAGAGAGAG 780
 DB 1089 AACTTTACCCCTCCAGATTTCCCAACAGCTCTTCTGTGAATCAACAAGTACAGAGAGAG 1148
 QY 781 GAGTGGCTACACAAACAGCTGAGGACGAAAGAGCAATGAATATCTGACATAGCTGAATTT 840
 DB 1149 GAGTGGCTACACAAACAGCTGAGGACGAAAGAGCAATGAATATCTGACATAGCTGAATTT 1208
 QY 841 TCGATTTTAAAGAAAGAAAGAAAGCAACCCAGATGGTTGAAGGATAAAGGAAACAAATTTG 900
 DB 1209 TCGATTTTAAAGAAAGAAAGAAAGCAACCCAGATGGTTGAAGGATAAAGGAAACAAATTTG 1268
 QY 901 TTTCGACGGGAAACTATTTTCGACAGTATCAATGATATATTTAGCCATAGAGCTTAAT 960
 DB 1269 TTTCGACGGGAAACTATTTTCGACAGTATCAATGATATATTTAGCCATAGAGCTTAAT 1328
 QY 961 AATAAGATGCCATATTTGATTTTGAACCGGGCTGCTTCCACCTAAAACTAAAAAACTTTA 1020
 DB 1329 AATAAGATGCCATATTTGATTTTGAACCGGGCTGCTTCCACCTAAAACTAAAAAACTTTA 1388
 QY 1021 CACAAGGCTATTTGAAGATTTCTTAAAGGCACTGGAAATTTATGATGCCACCTGTTACAGAC 1080
 DB 1389 CACAAGGCTATTTGAAGATTTCTTAAAGGCACTGGAAATTTATGATGCCACCTGTTACAGAC 1448
 QY 1081 AATGCTAATCAAGAAATGAAGGACATGTACGCTGCAAGCAGCATTTCTGCAACTAGAA 1140
 DB 1449 AATGCTAATCAAGAAATGAAGGACATGTACGCTGCAAGCAGCATTTCTGCTCACTAGAA 1508
 QY 1141 TTGTATGTAGAGGCTTACAGGATTTATGAAGCGGCACCTTAAGATTTGATCCATCCAAACAA 1200
 DB 1509 TTGTATGTAGAGGCTTACAGGATTTATGAAGCGGCACCTTAAGATTTGATCCATCCAAACAA 1568
 QY 1201 ATTGTACAAATTTGATGCTGAGAGATTCCGGAATGTAAATTTCAAGGAAACAGAACTAAATCT 1260
 DB 1569 ATTGTACAAATTTGATGCTGAGAGATTCCGGAATGTAAATTTCAAGGAAACAGAACTAAATCT 1628
 QY 1261 TAA 1263
 DB 1629 TAA 1631

RESULT 3
 ADB16939
 ID ADB16939 standard; cDNA; 1263 BP.
 XX
 AC ADB16939;
 XX
 DT 20-NOV-2003 (first entry)
 XX
 DE Pygmy chimpanzee DYXC1 cDNA sequence.
 XX
 KW gene; ss; pygmy chimpanzee; DYXC1; dyslexia; neurological disorder;
 KW reading disability; phonological processing; rapid naming;
 XX verbal short-term memory.
 XX Pan paniscus.
 XX

FH Key Location/Qualifiers
 FT CDS 1..1263
 FT /*tag= a
 FT /product= "DYX1 protein"
 XX WO2003068814-A1.
 XX
 XX
 XX
 XX PD 21-AUG-2003.
 XX
 XX 12-FEB-2003; 2003WO-FI000110.
 XX
 XX 12-FEB-2002; 2002US-0355782P.
 XX
 XX (LICN) LICENTIA LTD.
 XX
 XX Kere J, Taipale M, Nopola-Hemmi J, Kaminen N;
 XX WPI; 2003-646482/61.
 XX P-PSDB; ADB16940.
 DR
 DR
 XX New isolated, purified DYX1 nucleic acid for studying brain processes,
 PT e.g. reading, phonological processing, rapid naming or verbal short-term
 PT memory, or for diagnosing dyslexia or assessing the predisposition to
 PT dyslexia.
 XX
 XX Claim 29; Page 126-128; 135pp; English.
 PS
 XX This invention relates to a novel isolated human gene DYX1 that is
 CC functionally related to dyslexia, more particularly it describes single
 CC nucleotide polymorphisms thought to predispose an individual in to
 CC developing dyslexia. This is a neurological disorder with a genetic basis
 CC (DYX1 has been isolated to chromosome 15q21), which manifests itself as
 CC a specific reading disability. Specifically, DYX1 is can be useful in
 CC study of brain processes such as reading, phonological processing, rapid
 CC naming and verbal short-term memory. Accordingly, the present invention
 CC describes methods and materials for analysing allelic variations in the
 CC DYX1 gene, and also provides DYX1 as an antigen for the production of
 CC antibodies used in the diagnosis of dyslexia. This polynucleotide
 CC sequence is the pygmy chimpanzee DYX1 cDNA homologous to the human DYX1
 CC gene of the invention.
 XX
 SQ Sequence 1263 BP; 493 A; 213 C; 274 G; 283 T; 0 U; 0 Other;
 Query Match 99.2%; Score 1253.4; DB 9; Length 1263;
 Best Local Similarity 99.5%; Pred. No. 1.5e-261;
 Matches 1257; Conservative 0; Mismatches 6; Indels 0; Gaps 0;
 1 ATGCTCTTTCAGGTTAGCGATTACAGCTGGCGAGCAGACGAGCTGCGGTCTTTCTGCT 60
 1 ATGCCCCCTTCAGGTTAGCGATTACAGCTGGCGAGCAGACGAGCTGCGGTCTTTCTGCT 60
 61 CTGCCCCCTCAAGCGGTGTCGTGTCAGACACGCGACGTGTTCTGACGGAAACTATCTG 120
 61 CTGCCCCCTCAAGCGGTGTCGTGTCAGACACGCGACGTGTTCTGACGGAAACTATCTG 120
 121 AAGTCAACTTCTCCATTTTATTTAGGCAATTTCTGATGATGATGATGATGATGATGAT 180
 121 AAGTCAACTTCTCCATTTTATTTAGGCAATTTCTGATGATGATGATGATGATGATGAT 180
 181 AGCAGCAAGCAAGATTGGGAATGACACCATTTCTTCACTTGTATATAAAGAGAGCG 240
 181 AGCAGCAAGCAAGATTGGGAATGACACCATTTCTTCACTTGTATATAAAGAGAGCG 240
 241 GCCATGTGGAGACCCCTTCTGTCGCGGTGTTGACAAAGAGATGATGCAAGAAATAGA 300
 241 GCCATGTGGAGACCCCTTCTGTCGCGGTGTTGACAAAGAGATGATGCAAGAAATAGA 300
 301 GAAAAATCTATTTTACAGCAACAGAGAGCAAAAGAGCTACAGAGCAAAAGCTGCA 360
 301 GAAAAATCTATTTTACAGCAACAGAGAGCAAAAGAGCTACAGAGCAAAAGCTGCA 360
 361 GCAAAGCGGGAAGATCAAAATACGCTAAGTGTGATGATGATGATGATGATGATGATGAT 420

Db 361 GCAAAGCGGGAAGATCAAAATATGCACCTAAGTGTGATGATGATGATGATGATGATGATGAT 420
 Qy 421 AGGAAAAATATAGAGATATGAAGAAATGAACGATTAAGCCATTAAGCATTCGAA 480
 Db 421 AGGAAAAATATAGAGATATGAAGAAATGAACGATTAAGCCATTAAGCATTCGAA 480
 Qy 481 GCCTGGAAGAAATATCAAGAAAGCTGAGGAGCAAAAAAATTCAGAGAGAGAGAAA 540
 Db 481 GCCTGGAAGAAATATCAAGAAAGCTGAGGAGCAAAAAAATTCAGAGAGAGAGAAA 540
 Qy 541 TTATGTCAAAAGAAAGCAAAATTTAAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 600
 Db 541 TTATGTCAAAAGAAAGCAAAATTTAAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 600
 Qy 601 ACTAGAAATTTGGCATCTAGAAATCTTCTCCAAAGGGAGAAATTCAGAGAGAGAGAGAG 660
 Db 601 ACTAGAAATTTGGCATCTAGAAATCTTCTCCAAAGGGAGAGAAATTCAGAGAGAGAGAGAG 660
 Qy 661 ACTGAGAAAGTTAAAGGAG 720
 Db 661 ACTGAGAAAGTTAAAGGAG 720
 Qy 721 AACTTTACCCCTCGAGTATTTCCCAACAGCTCTTCCGTAATCAACAAGTAGCAGAGAGAG 780
 Db 721 AACTTTACCCCTCGAGTATTTCCCAACAGCTCTTCCGTAATCAACAAGTAGCAGAGAGAG 780
 Qy 781 GAGTGGCTACAAAACAGCTGAGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 840
 Db 781 GAGTGGCTGACAAAACAGCTGAGGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 840
 Qy 841 TGGATTTTAAAG 900
 Db 841 TGGATTTTAAAG 900
 Qy 901 TTTGCAACGAGAAACTATTTGGCAGCTATCAATGCATATTAATTTAGCCATAAGACTAAAT 960
 Db 901 TTTGCAACGAGAAACTATTTGGCAGCTATCAATGCATATTAATTTAGCCATAAGACTAAAT 960
 Qy 961 AATAAGATGACATTTGATTTTGAACCGGCTGCTTGCACCTTAAACTTAAACTTAAACTTAA 1020
 Db 961 AATAAGATGACATTTGATTTTGAACCGGCTGCTTGCACCTTAAACTTAAACTTAAACTTAA 1020
 Qy 1021 CACAAGGCTATTGAAGATTTCTTAAGGCACTGGAATTTATGATGCCACCTGTTTACAGAC 1080
 Db 1021 CACAAGGCTATTGAAGATTTCTTAAGGCACTGGAATTTATGATGCCACCTGTTTACAGAC 1080
 Qy 1081 AATGCTAATCAAGAAATGAAGGACATGTACGACGTGGAACAGCATTTCTGTCAACTAGAA 1140
 Db 1081 AATGCTAATCAAGAAATGAAGGACATGTACGACGTGGAACAGCATTTCTGTCAACTAGAA 1140
 Qy 1141 TTGTATGTAGAGCCCTACAGGATTTATGAAGCGGCACTTAAGATTGATCCATCCAAAGAA 1200
 Db 1141 TTGTATGTAGAGCCCTACAGGATTTATGAAGCGGCACTTAAGATTGATCCATCCAAAGAA 1200
 Qy 1201 ATTGTACAAATTCATGCTGAGAGAGATTCGGAATGTAAATTCAGAGAGAGAGAGAGAGAG 1260
 Db 1201 ATTGTACAAATTCATGCTGAGAGAGATTCGGAATGTAAATTCAGAGAGAGAGAGAGAGAG 1260
 Qy 1261 TAA 1263
 Db 1261 TAA 1263
 RESULT 4
 ADB16933
 ID ADB16933 standard; cDNA; 1263 BP.
 XX
 AC ADB16933;
 XX
 DT 20-NOV-2003 (first entry)
 XX
 DE Chimpanzee DYX1 cDNA sequence.
 XX

RESULT 6

ADB16937

ID ADB16937 standard; cDNA; 1263 BP.

XX

AC ADB16937;

XX

DT 20-NOV-2003 (first entry)

XX

DE Orangutan DYXC1 cDNA sequence.

XX

KW gene; ss; orangutan; DYXC1; dyslexia; neurological disorder;

KW reading disability; phonological processing; rapid naming;

KW verbal short-term memory.

XX

OS Pongo pygmaeus.

XX

FH Key Location/Qualifiers

FT CDS 1..1263

FT /*tag= a

FT /product= "DYXC1 protein"

XX

PN WO2003068814-A1.

XX

PD 21-AUG-2003.

XX

PF 12-FEB-2003; 2003WO-FI000110.

XX

PR 12-FEB-2002; 2002US-0355782P.

XX

PA (LICN) LICENTIA LTD.

XX

PI Kere J, Taipale M, Nopola-Hemmi J, Kaminen N;

XX

PI WPI; 2003-646482/61.

XX

DR P-P8DB; ADB16938.

XX

PT New isolated, purified DYXC1 nucleic acid for studying brain processes,

PT e.g. reading, phonological processing, rapid naming or verbal short-term

PT memory, or for diagnosing dyslexia or assessing the predisposition to

PT dyslexia.

XX

PS Claim 29; Page 123-124; 135pp; English.

XX

CC This invention relates to a novel isolated human gene DYXC1 that is functionally related to dyslexia, more particularly it describes single nucleotide polymorphisms thought to predispose an individual in to developing dyslexia. This is a neurological disorder with a genetic basis (DYXC1 has been isolated to chromosome 15q21), which manifests itself as a specific reading disability. Specifically, DYXC1 is can be useful in study of brain processes such as reading, phonological processing, rapid naming and verbal short-term memory. Accordingly, the present invention describes methods and materials for analysing allelic variations in the DYXC1 gene, and also provides DYXC1 as an antigen for the production of antibodies used in the diagnosis of dyslexia. This polynucleotide sequence is the orangutan DYXC1 cDNA homologous to the human DYXC1 gene of the invention.

XX

SQ Sequence 1263 BP; 493 A; 218 C; 272 G; 280 T; 0 U; 0 Other;

XX

Query Match

Best Local Similarity 98.5%; Score 1243.8; DB 9; Length 1263;

Matches 1251; Conservative 0; Mismatches 12; Indels 0; Gaps 0;

Qy 1 ATGCCTCTTCAGGTAGCGATTACAGCTGGCAGCAGACGAGACTGCGGTCTTCTGCT 60

Db 1 ATGCCCCCTCAGGTAGCGATTACAGCTGGCAGCAGACGAGACTGCGGTCTTCTGCT 60

Qy 61 CTGCCCCCTCAAGGGCGTGTGGCTCAGAGACAGGACGCTGTCTGCACGGAAAACTATCTG 120

Db 61 CTGCCCCCTCAAGGGCGTGTGGCTCAGAGACAGGACGCTGTCTGCACGGAAAACTATCTG 120

Qy 121 AAGGTCAACTTCTCTCCATTTTATTGAGGCAATTTCTTTATGCTCCCATAGACGATGAG 180
 Db 121 AAGGTCAACTTCTCTCCATTTTATTGAGGCAATTTCTTTATGCTCCCATAGACGATGAG 180
 Qy 181 AGCAGCAAAAGCAAGATTGGGAATGACACCACTTGTCTTCACTTGTATATAAAAGAAAGCG 240
 Db 181 AGCAGCAAAAGCAAGATTGGGAATGACACCACTTGTCTTCACTTGTATATAAAAGAAAGCG 240
 Qy 241 GCCATGTGGGAGACCCCTTTTCTGTGACGGGTGTGTGACAAAGAGATGATGCAAGAAATTAGA 300
 Db 241 GCCATGTGGGAGACCCCTTTTCTGTGACGGGTGTGTGACAAAGAGACGATGCAAGAAATTAGA 300
 Qy 301 GAAAAATCTATTTTACAGCACAAGAGAGACCAAAAGAGCTACAGAGCAAAAAGCTGCA 360
 Db 301 GAAAAATCTATTTTACAGCACAAGAGAGACCAAAAGAGCTACAGAGCAAAAAGCTGCA 360
 Qy 361 GCAAAGCGGGAAGATCAAAAAATAGCACTAAGTGTGATGATGAAGATTGAAGAAAGAG 420
 Db 361 GCAAAGCGGGAAGATCAAAAAATAGCACTAAGTGTGATGATGAAGATTGAAGAAAGAG 420
 Qy 421 AGGAAAAAATAGAAAGATATGAAAGAAATGAAACGGATAAAAAGCCATTAAGCATTTGAA 480
 Db 421 AGGAAAAAATAGAAAGATATGAAAGAAATGAAACGGATAAAAAGCCATTAAGCATTTGAA 480
 Qy 481 GCCTGGAAGAAATATCAAGAAAAAGCTGAGAGCAAAAAAATTCAGAGAGAAAGAAA 540
 Db 481 GCCTGGAAGAAATATCAAGAAAAAGCTGAGAGCAAAAAAATTCAGAGAGAAAGAAA 540
 Qy 541 TTATGTCAAAAAAGAAAGCAAAATTAAGAGAAAGAAAGAAAAAATAAATAATAAGAGTCTT 600
 Db 541 TTATGTCAAAAAAGAAAGCAAAATTAAGAGAAAGAAAGAAAAAATAAATAATAAGAGTCTT 600
 Qy 601 ACTAGAAATTTGGCATCTAGAAATCTTGTCTCCAAAAGGGAGAAAATTCAGAAAAATATATTT 660
 Db 601 ACTAGAAATTTGGCATCTAGAAATCTTGTCTCCAAAAGGGAGAAAATTCAGAAAAATATATTT 660
 Qy 661 ACTGAGAAGTTAAAGGAGACAGTATTTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 720
 Db 661 ACTGAGAAGTTAAAGGAGACAGTATTTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 720
 Qy 721 AACTTTTACCCCTCGAGTATTTCCCAACAGCTCTTCTGTGAATCAACAGTACAGAGAGGAG 780
 Db 721 AACTTTTACCCCTCGAGTATTTCCCAACAGCTCTTCTGTGAATCAACAGTACAGAGAGGAG 780
 Qy 781 GAGTGGCTTACACAAACAGCTGAGGCGACGAGACCAATGAATACTGACATAGCTGAACCTT 840
 Db 781 GAGTGGCTTACACAAACAGCTGAGGCGACGAGACCAATGAATACTGACATAGCTGAACCTT 840
 Qy 841 TGGCATTTAAAGAAAGAAAGAAAGAACCCAGAAATGGTTGAAGGATAAAGGAAACAAATTTG 900
 Db 841 TGGCATTTAAAGAAAGAAAGAAAGAACCCAGAAATGGTTGAAGGATAAAGGAAACAAATTTG 900
 Qy 901 TTTGCAACGGAAAACTATTTTGGCAGCTATCAATGCATATAATTTAGCCATAAGACTAAAT 960
 Db 901 TTTGCAACGGAAAACTATTTTGGCAGCTATCAATGCATATAATTTAGCCATAAGACTAAAT 960
 Qy 961 AATAAGATGCCATATTTGTATTTGAACCGGGCTGCTTGGCCACCTAAAAAATAAAACTTTA 1020
 Db 961 AATAAGATGCCATATTTGTATTTGAACCGGGCTGCTTGGCCACCTAAAAAATAAAACTTTA 1020
 Qy 1021 CACAAGGCTATTGAAGATTCTTCTTAAGGCACTGGAATTTATGATGCCACCTGTTTACAGAC 1080
 Db 1021 CACAAGGCTATTGAAGATTCTTCTTAAGGCACTGGAATTTATGATGCCACCTGTTTACAGAC 1080
 Qy 1081 AATCTAATGCAAGAAATGAAGGCAATGTACGAGTGGAAACAGCATTTCTGCTCAACTAGAA 1140
 Db 1081 AATCTAATGCAAGAAATGAAGGCAATGTACGAGTGGAAACAGCATTTCTGCTCAACTAGAA 1140
 Qy 1141 TTGTATGTAGAAAGCCCTACAGGATTATGAAGCGGCACCTTAAGATTGTATTCACCAACAAA 1200
 Db 1141 TTGTATGTAGAAAGCCCTACAGGATTATGAAGCGGCACCTTAAGATTGTATTCACCAACAAA 1200
 Qy 1201 ATTGTACAAATTGATGCTGAGAAAGATTTCGGGAATGTAAATTCAGGAACAGAACTAAATCT 1260

Db 1201 ATTGTACAAATTGATGCTGAGAGATTTCGGAAATGTAATTCAAGGACAGAACTAAAATCT 1260
Qy 1261 TAA 1263
Db 1261 TAA 1263
RESULT 7
ADC30210
ID ADC30210 standard; cDNA; 1641 BP.
XX
XX ADC30210;
AC
XX 18-DEC-2003 (first entry)
XX
XX Human novel cDNA sequence, SEQ ID NO:292.
XX
XX Human; diagnostic; drug screening; forensics; gene mapping;
KW biodiversity assessment; Parkinson's disease; Alzheimer's disease;
KW neurodegenerative diseases; anaemia; platelet disorder; wound; burns;
KW ulcers; osteoporosis; autoimmune disease; cancer;
KW molecular weight marker; food supplement; antiparkinsonian; nootropic;
KW neuroprotective; antianaemic; anticoagulant; thrombolytic; vulnerary;
KW antiulcer; osteopathic; immunosuppressive; antiinflammatory; cytostatic;
KW gene therapy; chromosome 15q21.3; gene; ss.
XX
XX Homo sapiens.
OS
XX WO2003029271-A2.
XX
XX 10-APR-2003.
XX
XX 24-SEP-2002; 2002WO-US030474.
XX
XX 24-SEP-2001; 2001US-0324631P.
XX
XX (HYSE-) HYSEQ INC.
XX
XX Tang TY, Zhang J, Ren F, Xue AJ, Zhao QA, Wang J, Wehrman T;
PI Zhou P, Ghosh M, Wang D, Ma Y, Aundi V, Wang Z, Weng G;
PI Haley-Vicente D, Drmanac RT;
XX
XX WPI; 2003-371981/35.
DR P-PSDB; ADC31181.
XX
XX New polynucleotide and polypeptide useful for diagnosing, preventing or
PT treating conditions such as neurodegenerative diseases, anemias, platelet
PT disorders, wounds, burns, ulcers, osteoporosis, autoimmune diseases or
PT cancer.
XX
XX Claim 1; SEQ ID NO 292; 1185pp; English.
XX
XX The invention relates to 971 novel human cDNA sequences (ADC29919-
CC ADC30899) and the polypeptides they encode (ADC30890-ADC31860). The
CC invention also relates to nucleic acid sequences over 99% identical with
CC the novel human cDNAs. The invention additionally encompasses expression
CC vectors and host cells comprising a nucleic acid of the invention; the
CC recombinant production of a polypeptide of the invention; an antibody
CC against a polypeptide of the invention; a method of detecting
CC polynucleotides or polypeptides of the invention; and methods of
CC identifying a compound which binds to a polypeptide of the invention. The
CC invention further discloses methods of preventing, treating or
CC ameliorating a medical condition; kits comprising polynucleotide probes
CC and/or monoclonal antibodies for carrying out the methods of the
CC invention; methods for the identification of compounds that modulate the
CC expression or activity of the polynucleotide and/or polypeptide; and 767
CC contig sequences corresponding to the cDNA sequences of the invention
CC (ADC31861-ADC32627) and the polypeptides encoded by the contigs (ADC32628
CC -ADC33394). The nucleic acids and polypeptides of the invention are
CC useful in diagnostics, drug screening, forensics, gene mapping, in the
CC identification of mutations responsible for genetic disorders or other
CC traits, for assessing biodiversity, and in producing many other types of

CC data and products dependent on DNA and amino acid sequences. They are
CC also used for treating diseases such as Parkinson's disease, Alzheimer's
CC disease and other neurodegenerative diseases, anaemia, platelet
CC disorders, wounds, burns, ulcers, osteoporosis, autoimmune diseases or
CC cancer. The nucleic acids may also be used as hybridisation probes or
CC primers, and in the recombinant production of a protein. The polypeptides
CC are also useful in generating antibodies, as molecular weight markers,
CC and as food supplements. The present sequence represents a specifically
CC claimed human cDNA sequence of the invention. Note: The sequence data for
CC this patent did not form part of the printed specification, but was
CC obtained in electronic format directly from WIPO at
CC ftp.wipo.int/pub/published_pct_sequences.
XX
XX Sequence 1641 BP; 614 A; 272 C; 331 G; 424 T; 0 U; 0 Other;
SQ
Query Match 83.1%; Score 1049; DB 10; Length 1641;
Best Local Similarity 100.0%; Pred. No. 2.9e-217;
Matches 1049; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 1 ATGCCTCTTCAGGTAGCGATTACAGCTGGCAGCAGACGAAGACTGGGTCCTTCTCTCT 60
Db 104 ATGCCTCTTCAGGTAGCGATTACAGCTGGCAGCAGACGAAGACTGGGTCCTTCTCTCT 163
Qy 61 CTGCCCCCTCAAAGCGGTGTCGTCAGAGACACGACGCTGTTCTCCACGAAAACTATCTG 120
Db 164 CTGCCCCCTCAAAGCGGTGTCGTCAGAGACACGACGCTGTTCTCCACGAAAACTATCTG 223
Qy 121 AAGTCAACTTTCTCTCAATTTTATTGAGGCAATTTCTTATGCTCCCATAGAGATGAG 180
Db 224 AAGTCAACTTTCTCTCAATTTTATTGAGGCAATTTCTTATGCTCCCATAGAGATGAG 283
Qy 181 AGCAGCAAAAGCAAGATTGGGAATGACACCATTTGTTTCACTTTGTATAAAAAAGAGCG 240
Db 284 AGCAGCAAAAGCAAGATTGGGAATGACACCATTTGTTTCACTTTGTATAAAAAAGAGCG 343
Qy 241 GCCATGTGGGAGACCCCTTTCTGTGACGGGTGTTGACAAAGAGATGATGCAAGAATTAGA 300
Db 344 GCCATGTGGGAGACCCCTTTCTGTGACGGGTGTTGACAAAGAGATGATGCAAGAATTAGA 403
Qy 301 GAAAAATCTATTTTACAAGCACAAGAGAGAGCAAAAGAGCTACAGAACGAAAAGCTGCA 360
Db 404 GAAAAATCTATTTTACAAGCACAAGAGAGAGCAAAAGAGCTACAGAACGAAAAGCTGCA 463
Qy 361 GCAAGCGGGAAGATCAAAAATAGCACAATAGTGTCTCATGATGAAGATTGAAGAGAAGAG 420
Db 464 GCAAGCGGGAAGATCAAAAATAGCACAATAGTGTCTCATGATGAAGATTGAAGAGAAGAG 523
Qy 421 AGGAAAAAATAGAGATATGAAAGAAAATGAAACGGATATAAAGCCACTAAAGCATTCGAA 480
Db 524 AGGAAAAAATAGAGATATGAAAGAAAATGAAACGGATATAAAGCCACTAAAGCATTCGAA 583
Qy 481 GCCTGGAAGAATATCAAGAAAAGCTGAGGAGCAAAAAAATTCAGAGAGAGAGAAA 540
Db 584 GCCTGGAAGAATATCAAGAAAAGCTGAGGAGCAAAAAAATTCAGAGAGAGAGAAA 643
Qy 541 TTATGTCAAAAGAAAAGCAAAATTAAGAAGAGAAAAGAAAATAAATAATAAGAGTCCTT 600
Db 644 TTATGTCAAAAGAAAAGCAAAATTAAGAAGAGAAAAGAAAATAAATAATAAGAGTCCTT 703
Qy 601 ACTAGAAAATTTGGCATCTAGAAAATCTTGTCTCCAAAAGGGAGAAAATTCAGAAAATATATTT 660
Db 704 ACTAGAAAATTTGGCATCTAGAAAATCTTGTCTCCAAAAGGGAGAAAATTCAGAAAATATATTT 763
Qy 661 ACTGAGAGATTAAAGGAAGACAGTATTCCTGCTCTCTGCTCTGTCAGTATTAAATC 720
Db 764 ACTGAGAGATTAAAGGAAGACAGTATTCCTGCTCTCTGTCAGTATTAAATC 823
Qy 721 AACTTTTCCCTCCAGTATTCCTGCTCTCTGCTCTGTCAGTATTAAATC 780
Db 824 AACTTTTCCCTCCAGTATTCCTGCTCTCTGCTCTGTCAGTATTAAATC 883
Qy 781 GAGTGGCTACACAAACAGCTGAGGCGACGAAGCAATGAATCTGACATAGCTGAACCT 840

Db 884 GAGTGGCTACACAAACAGCTGAGCCACGAAGAGCAATGAATCTGACATAGCTGAACCTT 943
Qy 841 TGCAGATTTAAAGAAAGAAAGAACCCAGAAATGGTTGAAGGATAAAGGAAACAAATTTG 900
Db 944 TGCAGATTTAAAGAAAGAAAGAAACCCAGAAATGGTTGAAGGATAAAGGAAACAAATTTG 1003
Qy 901 TTTCGAACGGAAACTATTTCGACGCTATCATGCTATATATTTAGCCATAGACTAAAT 960
Db 1004 TTTCGAACGGAAACCTATTTCGACGCTATCATGCTATATATTTAGCCATAGACTAAAT 1063
Qy 961 AATAAGATGCCACTATTGTTGTAATTTGAACCGGGCTGCTTGCACCTTAAACTTAAACCTTA 1020
Db 1064 AATAAGATGCCACTATTGTTGTAATTTGAACCGGGCTGCTTGCACCTTAAACTTAAACCTTA 1123
Qy 1021 CACAAGGCTATTGAAGATCTCTTAAGGC 1049
Db 1124 CACAAGGCTATTGAAGATCTCTTAAGGC 1152

RESULT 8
ID ADM01890 standard; cDNA; 1559 BP.
XX
AC ADM01890;
XX
DT 20-MAY-2004 (first entry)
XX
DE Human cDNA of the invention SEQ ID NO:575.
XX
KW ss; gene; human; gene therapy; diagnostic marker; pharmaceutical.
XX
OS Homo sapiens.
XX
FN EP1347046-A1.
XX
PD 24-SEP-2003.
XX
PF 12-APR-2002; 2002EP-00008400.
XX
PR 22-MAR-2002; 2002JP-00137785.
XX
PA (REAS-) RES ASSOC BIOTECHNOLOGY.
XX

PI Isogai T, Sugiyama T, Otsuki T, Wakamatsu A, Sato H, Ishii S;
PI Yanamoto J, Isono Y, Hio Y, Otsuka K, Nagai K, Irie R, Tamechika I;
PI Seki N, Yoshikawa T, Otsuka M, Nagahari K, Masuho Y;
XX
XX WPI; 2003-723558/69.
DR P-FSDB; ADM04333.
DR
XX
XX New polynucleotides and polypeptides are useful in gene therapy, for
XX developing a diagnostic marker or medicines for regulating their
XX expression and activity, or as a target of gene therapy.
XX
XX Claim 1; SEQ ID NO 575; 305pp; English.
XX

XX The invention relates to a novel human polynucleotide and the encoded
XX polypeptide. A polynucleotide of the invention may have a use in gene
XX therapy. An oligonucleotide of the invention ADM06202-ADM06773 is useful
XX as a primer for synthesizing the polynucleotide or as a probe for
XX detecting the polynucleotide. The polynucleotides ADM0316-ADM03758 are
XX useful in gene therapy, for developing a diagnostic marker or medicines
XX for regulating their expression and activity, or as a target of gene
XX therapy. The proteins ADM03759-ADM06201 encoded by the polynucleotides
XX are useful as pharmaceutical agents. The present sequence represents a
XX cDNA sequence of the invention.
XX
SQ Sequence 1559 BP; 592 A; 252 C; 309 G; 406 T; 0 U; 0 Other;

Query Match 82.9%; Score 1047.4; DB 11; Length 1559;
Best Local Similarity 99.9%; Pred. No. 6.4e-217;
Matches 1048; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 ATGCCTCTTCAGGTTAGCGATTACAGCTGGCAGCAGACGAGACTGCGGTCTTTCTGTCT 60
Db 43 ATGCCTCTTCAGGTTAGCGATTACAGCTGGCAGCAGACGAGACTGCGGTCTTTCTGTCT 102
Qy 61 CTGCCCCCTCAAAGCGGTGTGGTCTCAGAGACACGACGCTGTTCTGACGGAAGAACTATCTG 120
Db 103 CTGCCCCCTCAAAGCGGTGTGGTCTCAGAGACACGACGCTGTTCTGACGGAAGAACTATCTG 162
Qy 121 AAGTCAACTTTTCTCCATTTTATTTAGGCACTTTCTTTATGCTCCCATAGACGATGAG 180
Db 163 AAGTCAACTTTTCTCCATTTTATTTAGGCACTTTCTTTATGCTCCCATAGACGATGAG 222
Qy 181 AGCAGCAAGCAAGATTTGGGAATGACACCACTTCTTCCCTTGTATATAAAGAAAGCG 240
Db 223 AGCAGCAAGCAAGATTTGGGAATGACACCACTTCTTCCCTTGTATATAAAGAAAGCG 282
Qy 241 GCCATGTGGGAGACCCCTTTCTGTCAGCGGTGTGCAAAAGAGATGATGCAAGAAATAGA 300
Db 283 GCCATGTGGGAGACCCCTTTCTGTCAGCGGTGTGCAAAAGAGATGATGCAAGAAATAGA 342
Qy 301 GAAAAATCTATTTTACAAGCAAAAGAGAGACAAAGAGCTACAGAGCAAAAGCTGCA 360
Db 343 GAAAAATCTATTTTACAAGCAAAAGAGAGACAAAGAGCTACAGAGCAAAAGCTGCA 402
Qy 361 GCAAGCGGAGAGATCAAAAATACGCACTAAGTGTCTGATGAAGATTTGAAGAAAGAG 420
Db 403 GCAAGCGGAGAGATCAAAAATACGCACTAAGTGTCTGATGAAGATTTGAAGAAAGAG 462
Qy 421 AGGAAAAATATAGAAATATGAAAGAAATGAAAGGAGCAAAAGCTTAAAGCAATTTGAA 480
Db 463 AGGAAAAATATAGAAATATGAAAGAAATGAAAGGAGCAAAAGCTTAAAGCAATTTGAA 522
Qy 481 GCCTGGAAGAAATATCAAGAAAGAGTGGAGGACAAAGAAATTTTCAGAGAGAGAGAAA 540
Db 523 GCCTGGAAGAAATATCAAGAAAGAGTGGAGGACAAAGAAATTTTCAGAGAGAGAGAAA 582
Qy 541 TTATGTCAAAAGAAAGCAAAATTTAAAGAGGAGAAAGAAATATAAATATAAGAGCTTT 600
Db 583 TTATGTCAAAAGAAAGCAAAATTTAAAGAGGAGAAAGAAATATAAATATAAGAGCTTT 642
Qy 601 ACTAGAAATTTGGCATCTAGAAATCTTGTCTCAAAAGGAGAAATTCAGAAAAATATATT 660
Db 643 ACTAGAAATTTGGCATCTAGAAATCTTGTCTCAAAAGGAGAAATTCAGAAAAATATATT 702
Qy 661 ACTGAGAAGTTAAAGGAGAGACAGTATTCCTGCTCCTCTGTTGGCAGTATTAAATC 720
Db 703 ACTGAGAAGTTAAAGGAGAGACAGTATTCCTGCTCCTCTGTTGGCAGTATTAAATC 762
Qy 721 AACTTTACCTCGAGTATTTCCCAAGAGCTTCTGTAATCACAAGTAGCAGAGAGAG 780
Db 763 AACTTTACCTCGAGTATTTCCCAAGAGCTTCTGTAATCACAAGTAGCAGAGAGAG 822
Qy 781 GAGTGGCTACACAAACAGCTGAGGACGAGAGCAATGAATCTGACATAGCTGAACCTT 840
Db 823 GAGTGGCTACACAAACAGCTGAGGACGAGAGCAATGAATCTGACATAGCTGAACCTT 882
Qy 841 TGCAGATTTAAAGAAAGAAAGAAACCCAGAAATGGTTGAAGGATAAAGGAAACAAATTTG 900
Db 883 TGCAGATTTAAAGAAAGAAAGAAACCCAGAAATGGTTGAAGGATAAAGGAAACAAATTTG 942
Qy 901 TTTCGAACGGAAACTATTTCGACGCTATCATGCTATATATTTAGCCATAGACTAAAT 960
Db 943 TTTCGAACGGAAACTATTTCGACGCTATCATGCTATATATTTAGCCATAGACTAAAT 1002
Qy 961 AATAAGATGCCACTATTGTTGTAATTTGAACCGGGCTGCTTGCACCTTAAACTTAAACCTTA 1020
Db 1003 AATAAGATGCCACTATTGTTGTAATTTGAACCGGGCTGCTTGCACCTTAAACTTAAACCTTA 1062
Qy 1021 CACAAGGCTATTGAAGATCTCTTAAGGC 1049
Db 1063 CACAAGGCTATTGAAGATCTCTTAAGGC 1091


```
RESULT 9
ADB16924
ID ADB16924 standard; cDNA; 1697 BP.
XX
AC ADB16924;
XX
20-NOV-2003 (first entry)
XX
cDNA sequence of the murine DYXC1 mRNA.
XX
gene; ss; mouse; DYXC1; dyslexia; neurological disorder;
XX reading disability; phonological processing; rapid naming;
XX verbal short-term memory; murine.
XX
OS Mus musculus.
XX
FH Key Location/Qualifiers
FT CDS 48..1310
FT /tag= a
FT /product= "DYXC1 protein"
XX
WO2003068814-A1.
XX
21-AUG-2003.
XX
12-FEB-2003; 2003WO-FI000110.
XX
12-FEB-2002; 2002US-0355782P.
XX
(LICN ) LICENTIA LTD.
XX
Kere J, Taipale M, Nopola-Hemmi J, Kaminen N;
XX
WPI; 2003-646482/61.
DR P-PSDB; ADB16925.
XX
New isolated, purified DYXC1 nucleic acid for studying brain processes,
XX e.g. reading, phonological processing, rapid naming or verbal short-term
XX memory, or for diagnosing dyslexia or assessing the predisposition to
XX dyslexia.
XX
Disclosure; Page 51-53; 135pp; English.
XX
This invention relates to a novel isolated human gene DYXC1 that is
XX functionally related to dyslexia, more particularly it describes single
XX nucleotide polymorphisms thought to predispose an individual in to
XX developing dyslexia. This is a neurological disorder with a genetic basis
XX (DYXC1 has been isolated to chromosome 15q21), which manifests itself as
XX a specific reading disability. Specifically, DYXC1 is can be useful in
XX study of brain processes such as reading, phonological processing, rapid
XX naming and verbal short-term memory. Accordingly, the present invention
XX describes methods and materials for analysing allelic variations in the
XX DYXC1 gene, and also provides DYXC1 as an antigen for the production of
XX antibodies used in the diagnosis of dyslexia. This polynucleotide is the
XX cDNA sequence of the murine DYXC1 mRNA of the invention.
XX
SQ Sequence 1697 BP; 534 A; 360 C; 441 G; 362 T; 0 U; 0 Other;
Query Match 66.6%; Score 840.8; DB 9; Length 1697;
Best Local Similarity 80.3%; Pred. No. 3.5e-172;
Matches 1012; Conservative 0; Mismatches 242; Indels 6; Gaps 2;
Qy 1 ATGCCTCTTACAGTTAGCGATTACAGCTGGCAGCAGCAGAACTGGGCTTCTTGTCT 60
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
48 ATGCCAGTGCAGTGCAGGAATTCAGCTGGCAGCAGCAGCGCGCAGCATCTCTCTGTCG 107
Qy 61 CTGCCCTCAAGCGCTGCTGCAGACACGACGCTGTTCTGCACGGAACCTATCTG 120
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
108 CTGCTCTGCGGGGCGTCTGCTGGCGCATCTGACGATTTCTGTGGGAAGTACCTG 167
Qy 121 AAGGTCACACTTCTCCATTTTATTTTGGAGCATTTCTTTATGTCTCCCATAGACATGAG 180
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
168 AAGGTTAACTTCTCTCCATTTTATTTTGGAGCTGTTTCTCTATGCTCTCCCATAGATGGG 227
```

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Qy 181 AGCAGCAAAAGCAAGATTGGGAATGACACCATTTGTCTTACCTTGTATATAAAGAAGCG 240
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
228 AAGAGCAAAAGCCAAAGATTGGAATGACACCATTTCTTTTTCATTTGATATAAAGAGGCCA 287
Qy 241 GCCATGTGGGAGACCCCTTTCTGTGACGGGTGTTGACAAAGAGATGATGCAAGAAATTAGA 300
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
288 GTTCTGTGGGATAGCCCTTTCTGTGCGGGTGTGATAAAGAGATGATGACAGAAATAGA 347
Qy 301 GAAAAATCTATTTTACAGCACAAGAGAGCAAAAGAGCTACAGAACCAAGCTGCA 360
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
348 GAAAAATCTATCTTTCAGCACAAGAGAAACCAAGAGCCACAGAAAGCAAAAGCTGTT 407
Qy 361 GCAAGCGGGAAGATCAAAATACGCATTAAGTGTATGATGAAGATTGAAAGAGAAGAG 420
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
408 GCCAAGCGAGAGACCCAGATACGCCTAGGCGAGTGTGAAGATTGAAAGAGAAGAG 467
Qy 421 AGGAAAAAATAGAGATATGAAAGAAAATGAACCGGATAAAAGCCACTAAAGCATTTGAA 480
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
468 AGGAAAAAATCTCGAAGATCTGAAAGAAAATGAACCGGAAAAGCAACTAGCGAATTAGA 527
Qy 481 GCCTGGAAGAATATCAAGAAAAGCTGAGGAGCAAAAAAATAAATTTAGAGAGAAGAAA 540
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
528 GCCTGGAAGAATGTCAAAAGAAAAGCTGACGCAAAAAAAGAGTCCAGAGGAAAGGAGAA 586
Qy 541 TTATGTCAAAAAGAAAAGCAAAATTAAGAGGAGGAAAAAATAAATAATTAAGAGTCTT 600
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
587 --ACCGCTCGAGGAAAGCAAGCT---GAAGAGACCAAGCTCTAAACCTCGGGGTTTG 641
Qy 601 ACTAGAAAATTTGGCATCTAGAAAATCTTGTCTCAAAAGGGGAGAAATTCAGAAAAATAT 660
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
642 CCCGGAAGGCCCCACCCTCCCTCCCAAGAGGAGGAGGAAATTCGGAAGAAACATATTT 701
Qy 661 ACTGAGAAGTTAAAGGAAGACAGTATTCCTGCTCTCTGCTGCTGTGTCAGTATTAAATC 720
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
702 CCTGAGAAGTTAAAGGAAGACAGAGTCCCTGCGCTCGCTCCCTGCGAGTATTCAAAATC 761
Qy 721 AACTTTACCCCTCGAGTATTCCCAACAGCTCTTCGTGAATCAACAAGTAGCAGAGAGGAG 780
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
762 AGCTTTACCCCTCGAGTATTCCCAACAGCTCTTCGCGAATCCCAAGTCCGCAAGAGGAG 821
Qy 781 GAGTGGCTACACAAACAGCTGAGGACGAGAGCAATGAATACTGTACATAGCTGAACCTT 840
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
822 GAGTGGCTGCATAAACAAGCAGAGACACGAGAGCCATGAGCACTGACCTTCTCTGAGTTC 881
Qy 841 TGGGATTTAAAGAGAGAAAGAACCCAGATGTTGAGGATTAAGAGAAACAAATTTG 900
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
882 TTTGACTTAAAGAGAGAGAGGAATCCAGACTGGTGTGAAGACAAAGGGAACAAATTTG 941
Qy 901 TTTGCAACGGAAACTATTTGGCAGCTATCAATGCATATAATTTAGCCATAAGACTAAAT 960
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
942 TTTGCAACAGAAACTATTTGGCAGCGGTTGATGATATTAATTTAGCCATACGACTGAC 1001
Qy 961 AATAAGATGCCACTATTGTTATTTGAACCGGGCTGCTGCCACCTTAAACCTAAAAA 1020
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1002 TGTAAAGATCCCATTTATTTGATTTGATTCGGGCTGCTTGCACCTCAAAATTA 1061
Qy 1021 CACAGGCTATTGAGATTTCTTAAAGCACTGGAATTTATGATGCCACCTGTTTACAGAC 1080
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1062 CACAAGGCCATTCGAGGACTCTTCTTAAGGCACTAGAGTTATTTGACACCACTGTTGCTGAC 1121
Qy 1081 AATGCTAATGCAAGATTAAGGACATGTACGCTGGAACAGCATTTCTGTCAACTAGAA 1140
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1122 AATGCCAATGCAAGATTAAGGACACACGTCGACGAGGACAGGCTTCTGTCACTAGAA 1181
Qy 1141 TTGTATGTAGAGGCCCTACAGGATTTAAGAGCGGCACTTAAGATTGATTCATCCAA 1200
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1182 TTGTATGTGAAGGCTTTCGAAGATTAAGAGCTGCACTTAAGATTGACCCAGCCACACA 1241
Qy 1201 ATTGTACAAATTTGATGCTGAGAGATTTCCGAAATGTAATTCAGGAAACAGACTAAATCT 1260
Dy ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
1242 GTTGTACAGAACGATCAGAGAGATTTCCGAAATATAATTCAGGGAGCGGCACTGAAGTCT 1301
```


RESULT 10
 ADC32116
 ID ADC32116 standard; cDNA; 608 BP.
 XX
 AC ADC32116;
 XX
 DT 18-DEC-2003 (first entry)
 XX
 DE Human novel cDNA contig sequence, SEQ ID NO:2198.
 XX
 KW Human; diagnostic; drug screening; forensics; gene mapping;
 KW biodiversity assessment; Parkinson's disease; Alzheimer's disease;
 KW neurodegenerative diseases; anaemia; platelet disorder; wound; burns;
 KW ulcers; osteoporosis; autoimmune disease; cancer;
 KW molecular weight marker; food supplement; antiparkinsonian; nootropic;
 KW neuroprotective; anti-anaemic; anticoagulant; thrombolytic; vulnerary;
 KW anticancer; osteopathic; immunosuppressive; antiinflammatory; cytostatic;
 KW gene therapy; chromosome 15q21.3; ss.
 XX
 OS Homo sapiens.
 XX
 XX WO2003029271-A2.
 XX
 PD 10-APR-2003.
 XX
 XX 24-SEP-2002; 2002WO-US030474.
 XX
 XX 24-SEP-2001; 2001US-0324631P.
 XX
 XX (HYSE-) HYSEQ INC.
 XX
 XX Tang TY, Zhang J, Ren F, Xue AJ, Zhao QH, Wang J, Wehrman T;
 PI Zhou P, Ghosh M, Wang D, Ma Y, Asundi V, Wang Z, Weng G;
 PI Haley-Vicente D, Drmanac RT;
 XX
 XX WPI; 2003-371981/35.
 DR P-PSDB; ADC32883.
 XX
 PT New polynucleotide and polypeptide useful for diagnosing, preventing or
 PT treating conditions such as neurodegenerative diseases, anemias, platelet
 PT disorders, wounds, burns, ulcers, osteoporosis, autoimmune diseases or
 PT cancer.
 XX
 XX Example 2; SEQ ID NO 2198; 1185pp; English.
 PS
 PS The invention relates to 971 novel human cDNA sequences (ADC29919-
 CC ADC30889) and the polypeptides they encode (ADC30890-ADC31860). The
 CC invention also relates to nucleic acid sequences over 99% identical with
 CC the novel human cDNAs. The invention additionally encompasses expression
 CC vectors and host cells comprising a nucleic acid of the invention; the
 CC recombinant production of a polypeptide of the invention; an antibody
 CC against a polypeptide of the invention; a method of detecting
 CC polynucleotides or polypeptides of the invention; and methods of
 CC identifying a compound which binds to a polypeptide of the invention. The
 CC invention further discloses methods of preventing, treating or
 CC ameliorating a medical condition; kits comprising polynucleotide probes
 CC and/or monoclonal antibodies for carrying out the methods of the
 CC invention; methods for the identification of compounds that modulate the
 CC expression or activity of the polynucleotide and/or polypeptide; and 767
 CC contig sequences corresponding to the cDNA sequences of the invention
 CC (ADC31861-ADC32627) and the polypeptides encoded by the contigs (ADC32628
 CC -ADC33394). The nucleic acids and polypeptides of the invention are
 CC useful in diagnostics, drug screening, forensics, gene mapping, in the
 CC identification of mutations responsible for genetic disorders or other
 CC traits, for assessing biodiversity, and in producing many other types of
 CC data and products dependent on DNA and amino acid sequences. They are
 CC also used for treating diseases such as Parkinson's disease, Alzheimer's
 CC disease and other neurodegenerative diseases, anaemia, platelet
 CC disorders, wounds, ulcers, osteoporosis, autoimmune diseases or
 CC cancer. The nucleic acids may also be used as hybridisation probes or
 CC primers, and in the recombinant production of a protein. The polypeptides
 CC are also useful in generating antibodies, as molecular weight markers,

CC and as food supplements. The present sequence represents a human contig
 CC sequence used in an example of the invention. Note: The sequence data for
 CC this patent did not form part of the printed specification, but was
 CC obtained in electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 608 BP; 230 A; 103 C; 135 G; 140 T; 0 U; 0 Other;

Query Match 41.5%; Score 524; DB 10; Length 608;
 Best Local Similarity 100.0%; Pred. No. 1.1e-103;
 Matches 524; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 524 TTCAGAGAGAGAGAAATTTATGTCAAAAGAAAAGCAAAATTAAGAAGAGAGAAAAAAA 583

Db 1 TTCAGAGAGAGAGAAATTTATGTCAAAAGAAAAGCAAAATTAAGAAGAGAGAAAAAAA 60

QY 584 TAAATAAAGAGCTTACTAGAAATTTGGCATCTAGAAATCTTGCTCCAAAAGGGAGAA 643

Db 61 TAAATAAAGAGCTTACTAGAAATTTGGCATCTAGAAATCTTGCTCCAAAAGGGAGAA 120

QY 644 ATTCAGAAAATATATTTACTGAGAAAGTTAAAGGAGACAGTATTCTCTCTCTCTCTG 703

Db 121 ATTCAGAAAATATATTTACTGAGAAAGTTAAAGGAGACAGTATTCTCTCTCTCTCTG 180

QY 704 TTGGCAGTATTAAATCAACTTTTACCCCTCGAGTATTTCCCAACAGCTTTCTGTGAATCAC 763

Db 181 TTGGCAGTATTAAATCAACTTTTACCCCTCGAGTATTTCCCAACAGCTTTCTGTGAATCAC 240

QY 764 AAGTAGCAGAGAGAGAGAGGAGTGCTACACAAACAGCTGAGGCAGAGAGCAATGAATA 823

Db 241 AAGTAGCAGAGAGAGAGAGGAGTGCTACACAAACAGCTGAGGCAGAGAGCAATGAATA 300

QY 824 CTGCATAGCTGAACTTTTGGCATTTAAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 883

Db 301 CTGCATAGCTGAACTTTTGGCATTTAAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 360

QY 884 ATAAAGGAG 943

Db 361 ATAAAGGAG 420

QY 944 TAGCCATAAG 1003

Db 421 TAGCCATAAG 480

QY 1004 TAAAACTAAAAAACTTACACAGGCTATTGGAAGATTCTTCTTAAG 1047

Db 481 TAAAACTAAAAAACTTACACAGGCTATTGGAAGATTCTTCTTAAG 524

RESULT 11

ACH35463

ID ACH35463 standard; cDNA; 488 BP.

XX ACH35463;

AC ACH35463;

DT 13-OCT-2003 (first entry)

DE Human endothelial cell cDNA #3596.

KW Human; ss; sequencing by hybridisation; SBH; expressed sequence tag; EST;

XX genome mapping; biodiversity; genetic disorder.

OS Homo sapiens.

XX US2003073623-A1.

PD 17-APR-2003.

XX 30-JUL-2001; 2001US-00918995.

PR 30-JUL-2001; 2001US-00918995.

XX (DRMA/) DRMANAC R T.

```
PA (LABA/) LABAT I.
PA (STAC/) STACHE-CRAIN B.
PA (DICK/) DICKSON M C.
PA (JONE/) JONES L W.
XX
XX Drmanac RT, Labat I, Stache-Crain B, Dickson MC, Jones LW;
XX
XX WPI; 2003-615964/58.
XX
XX New polynucleotide sequences obtained from various cDNA libraries, useful
XX as hybridization probes, as oligomers for PCR, for chromosome and gene
XX mapping, in the recombinant production of protein, or in generating
XX antisense DNA or RNA.
XX
XX Claim 1; SEQ ID NO 22675; 44pp; English.
XX
XX The invention relates to an isolated polynucleotide comprising any one of
XX 38043 cDNA sequences, appearing as ACH12789-ACH50831, whose sequence was
XX determined by the technique of SBH (sequencing by hybridisation). Also
XX included is a purified polypeptide comprising a sequence corresponding to
XX a reading frame of the novel polynucleotide. The nucleic acid sequences
XX are useful in diagnostics as expressed sequence tags (EST) for
XX identifying expressed genes or for physical mapping of the human genome,
XX in forensics, in assessing biodiversity, or in identifying mutations
XX responsible for genetic disorders and other traits. The nucleotide
XX sequences are also useful as hybridisation probes, as oligomers for PCR,
XX for chromosome and gene mapping, in the recombinant production of
XX protein, or in generating antisense DNA or RNA. The purified polypeptide
XX is useful for generating antibodies specific for it. The present sequence
XX is one of the 38043 isolated cDNA/EST sequences. Note: The sequence data
XX for this patent did not form part of the printed specification, but was
XX obtained in electronic format directly from USPTO at
XX seqdata.uspto.gov/sequence.html?DocID=20030073623
XX
XX Sequence 488 BP; 141 A; 105 C; 126 G; 116 T; 0 U; 0 Other;
XX
XX Query Match 29.3%; Score 369.8; DB 9; Length 488;
XX Best Local Similarity 99.5%; Pred. No. 2.4e-70;
XX Matches 371; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
XX
XX 35 AGACGAAGACTGCGGTCTTTCTGTCCTGCCCCCTCAAAGGCGTGTGCGTCAGACACGG 94
XX |
XX 28 AGACGAAGACTGCGGTCTTTCTGTCCTGCCCCCTCAAAGGCGTGTGCGTCAGACACGG 87
XX |
XX 95 ACGTGTCTGACCGGAAAACATCTCTGAAGTCACTTCCTCCATTTTATTGAGGCAT 154
XX |
XX 88 ACGTGTCTGACCGGAAAACATCTCTGAAGTCACTTCCTCCATTTTATTGAGGCAT 147
XX |
XX 155 TTCTTTATGCTCCATAGACGATGAGACGACGACGACGACGACGACGACGACGACGACG 214
XX |
XX 148 TTTCTTTATGCTCCATAGACGATGAGACGACGACGACGACGACGACGACGACGACGACG 207
XX |
XX 215 TTTTCACCTTGATATAAAAAAGAGCGGCGCATGTGGGAGACCCCTTTCTGTGACGGGTGTG 274
XX |
XX 208 TTTTCACCTTGATATAAAAAAGAGCGGCGCATGTGGGAGACCCCTTTCTGTGACGGGTGTG 267
XX |
XX 275 ACAAGAGATGATGCAAGAAATTAGAGAAAATCTATTATCAAGACCAAGAGAGACGAA 334
XX |
XX 268 ACAAGAGATGATGCAAGAAATTAGAGAAAATCTATTATCAAGACCAAGAGAGACGAA 327
XX |
XX 335 AAGAAGCTTACAGAGCAAAAGCTCAGCAAGAGCGGGAAGATCAAAAATACGCCTAAAGTG 394
XX |
XX 328 AAGAAGCTTACAGAGCAAAAGCTCAGCAAGAGCGGGAAGATCAAAAATACGCCTAAAGTG 387
XX |
XX 395 TCATGATGAAGAT 407
XX |
XX 388 TCATGATGAAGCT 400
XX |
XX
XX RESULT 12
XX ACH23091
XX ID ACH23091 standard; cDNA; 458 BP.
XX
XX AC ACH23091;
XX XX
XX XX 13-OCT-2003 (first entry)
XX DE
XX XX Human adult ovary cDNA #1471.
XX XX
XX XX Human; ss; sequencing by hybridisation; SBH; expressed sequence tag; EST;
XX KW genome mapping; biodiversity; genetic disorder.
XX XX
XX XX Homo sapiens.
XX XX
XX XX US2003073623-A1.
XX PN
XX XX 17-APR-2003.
XX PD
XX XX
XX PF 30-JUL-2001; 2001US-00918995.
XX PR
XX PR 30-JUL-2001; 2001US-00918995.
XX XX
XX PA (DRMA/) DRMANAC R T.
XX PA (LABA/) LABAT I.
XX PA (STAC/) STACHE-CRAIN B.
XX PA (DICK/) DICKSON M C.
XX PA (JONE/) JONES L W.
XX XX
XX PI Drmanac RT, Labat I, Stache-Crain B, Dickson MC, Jones LW;
XX XX
XX XX WPI; 2003-615964/58.
XX DR
XX XX New polynucleotide sequences obtained from various cDNA libraries, useful
XX as hybridization probes, as oligomers for PCR, for chromosome and gene
XX mapping, in the recombinant production of protein, or in generating
XX antisense DNA or RNA.
XX
XX Claim 1; SEQ ID NO 10303; 44pp; English.
XX
XX The invention relates to an isolated polynucleotide comprising any one of
XX 38043 cDNA sequences, appearing as ACH12789-ACH50831, whose sequence was
XX determined by the technique of SBH (sequencing by hybridisation). Also
XX included is a purified polypeptide comprising a sequence corresponding to
XX a reading frame of the novel polynucleotide. The nucleic acid sequences
XX are useful in diagnostics as expressed sequence tags (EST) for
XX identifying expressed genes or for physical mapping of the human genome,
XX in forensics, in assessing biodiversity, or in identifying mutations
XX responsible for genetic disorders and other traits. The nucleotide
XX sequences are also useful as hybridisation probes, as oligomers for PCR,
XX for chromosome and gene mapping, in the recombinant production of
XX protein, or in generating antisense DNA or RNA. The purified polypeptide
XX is useful for generating antibodies specific for it. The present sequence
XX is one of the 38043 isolated cDNA/EST sequences. Note: The sequence data
XX for this patent did not form part of the printed specification, but was
XX obtained in electronic format directly from USPTO at
XX seqdata.uspto.gov/sequence.html?DocID=20030073623
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XX Sequence 458 BP; 170 A; 86 C; 97 G; 102 T; 0 U; 3 Other;
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XX Query Match 26.6%; Score 336; DB 9; Length 458;
XX Best Local Similarity 91.1%; Pred. No. 5e-63;
XX Matches 368; Conservative 0; Mismatches 35; Indels 1; Gaps 1;
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XX 56 GAGAAAGACTGACATATATTTACTGAGACGTTAAAGAGAGACAGATATTCGACCTCT 115
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XX 699 CTCTGTTGGCAGTATTAAATCAACTTTACCCCTCGAGTATTCCCAACAGCTCTTCGTGA 758
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XX 116 GTGTGCATGGCGCTA-CCCCACCAACCCGACGTCGAGAAATTCCAATCAGTCTTCTCGA 174
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XX 819 GAATACCTGACATAGCTGAACTTTTTCGATTTTAAAGAGAGAGAAAGAACCCAGCAATGGTT 878
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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: July 8, 2005, 01:40:13 ; Search time 4597 Seconds
(without alignments)
10457.945 Million cell updates/sec

Title: US-10-681-199-1
Perfect score: 1263
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Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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2: gb_est2: *
3: gb_hc: *
4: gb_est3: *
5: gb_est4: *
6: gb_est5: *
7: gb_est6: *
8: gb_gss1: *
9: gb_gss2: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1034.8	81.9	1600	3	BC017392 Homo sapi
2	812.6	64.3	844	6	CD107587 AGENCOURT
3	676.4	53.6	933	6	CD358543 AGENCOURT
4	667.8	52.9	793	7	CO738958 SILE04c21
5	567.4	44.9	792	4	BG711796 602720472
6	553	43.8	788	2	BES64350 601343161
7	519	41.1	683	5	BP460416 BP460416
8	513.8	40.7	1190	3	AK005832 Mus muscu
9	510.4	40.4	1131	5	BQ217312 AGENCOURT
10	508	40.2	917	6	BY705981 BY705981
11	488.6	38.7	545	2	BF216970 601884034
12	474.4	37.6	535	7	CK478982 UI-CF-FNO
13	449	35.6	796	7	CK603982 AGENCOURT
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15	427	33.8	1183	2	BF207765 601861861
16	422	33.4	462	7	CN429788 170006000
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18	409.6	32.4	988	4	BI517373 603041624
19	408	32.3	559	5	BU607404 UI-CF-FNO
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23	385.8	30.5	651	5	BU852580 AGENCOURT
24	376.8	29.8	844	7	CK598370 AGENCOURT

c	25	374.4	29.6	772	6	CA422987	UI-H-FLO-
	26	371.8	29.4	665	4	BG540324	BG540324 602568825
	27	364	28.8	573	2	BE972748	BE972748 601652170
	28	363.2	28.8	775	4	BG206612	BG206612 RST26063
	29	350.8	27.8	477	6	CB545994	CB545994 AMGNNUC:M
	30	344.4	27.3	744	4	BG192162	BG192162 RST11269
	31	341	27.0	752	4	BG183140	BG183140 RST2158 A
c	32	325	25.7	325	1	A1783611	A1783611 t299601.x
	33	308.4	24.4	375	2	AW481240	AW481240 35107 MAR
	34	301.8	23.9	687	4	BG242087	BG242087 602354631
	35	298	23.6	764	5	BU307742	BU307742 603539279
	36	296.2	23.5	767	5	BX867086	BX867086 BX867086
	37	295	23.4	855	2	BF248143	BF248143 601859338
c	38	293.2	23.2	727	4	BJ056709	BJ056709 BJ056709
	39	291.8	23.1	478	2	BB615485	BB615485 BB615485
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	42	283.8	22.5	470	1	AV266346	AV266346 AV266346
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ALIGNMENTS

RESULT 1
BC017392
LOCUS BC017392 Homo sapiens, Similar to RIKEN cDNA 1700010124 gene, clone
DEFINITION IMAGE:4081622, mRNA.
ACCESSION BC017392
VERSION BC017392.1 GI:19263480
KEYWORDS HTC.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 1600)
AUTHORS Strausberg,R.
TITLE Direct Submission
JOURNAL Submitted (13-NOV-2001) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
USA
REMARK NIH-MGC Project URL: http://mgc.nci.nih.gov
COMMENT Contact: MGC help desk
Email: cgapbs-remail.nih.gov
Tissue Procurement: ATCC
CDNA Library Preparation: CLONTECH Laboratories, Inc.
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Institute for Systems Biology
http://www.systemsbio.org
contact: amadan@systemsbio.org
Anup Madan, Jessica Fahey, Erin Helton, Mark Kettelman, Anuradha
Madan, Stephanie Rodriguez, Amy Sanchez and Michelle Whiting
Clone distribution: MGC clone distribution information can be found
through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov
Series: IRAL Plate: 32 Row: k Column: 22
This clone was selected for full length sequencing because it
passed the following selection criteria: Hexamer frequency ORF
analysis, Genomescan gene prediction
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Location/Qualifiers
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/mol_type="mRNA"
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/note="Vector: pDNR-LIB"		ORIGIN	
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Db	53	ATGCCTCTTCAGGTTAGCGATTACAGCTGGCAGCAGACGAGACTGCGGTCTTCTGTCT	112
Qy	61	CTGCCCTCAAAGCGGTGTGCGTCAGAGACACGACGTGTTCTGCACGGAAACATCTCTG	120
Db	113	CTGCCCTCAAAGCGGTGTGCGTCAGAGACACGACGTGTTCTGCACGGAAACATCTCTG	172
Qy	121	AAGTCAACTTCTCCTCAATTTTATTTAGGCAATTCCTTATGTCTCCATAGACGATGAG	180
Db	173	AAGTCAACTTCTCCTCAATTTTATTTAGGCAATTCCTTATGTCTCCATAGACGATGAG	232
Qy	181	AGCAGCAACCAAGATTGGGAATGACACCAATTCCTTACCTTGTATATAAAGAAAGCG	240
Db	233	AGCAGCAACCAAGATTGGGAATGACACCAATTCCTTACCTTGTATATAAAGAAAGCG	292
Qy	241	GCCATGTGGGAGACCCCTTCTGTGACGGGTGTCACAAAGAGATGATGCAAGAAATTAGA	300
Db	293	GCCATGTGGGAGACCCCTTCTGTGACGGGTGTCACAAAGAGATGATGCAAGAAATTAGA	352
Qy	301	GAATAATCTATTTTACAAGCACAAGAGAGACAAAGAGCTACAGAGCAAAAGCTGCA	360
Db	353	GAATAATCTATTTTACAAGCACAAGAGAGACAAAGAGCTACAGAGCAAAAGCTGCA	412
Qy	361	GCAAGCGGGAAGATCAAAATACGCTAAGTGTCTCATGTGATGATGAGATTGAAGAAGAAG	420
Db	413	GCAAGCGGGAAGATCAAAATACGCTAAGTGTCTCATGTGATGAGATTGAAGAAGAAG	472
Qy	421	AGG-AAAAAATAGAAGATATGAAGAAATGAACGGATAAAGCCACTAAAGCATTTGGA	479
Db	473	AGG-AAAAAATAGAAGATATGAAGAAATGAACGGATAAAGCCACTAAAGCATTTGGA	532
Qy	480	AGCTGGAAGAATATCAAGAAAGCTGAGGAGCAAAAAAATTCAGAGAGAAGAGAA	539
Db	533	AGCTGGAAGAATATCAAGAAAGCTGAGGAGCAAAAAAATTCAGAGAGAAGAGAA	592
Qy	540	ATTATGTCAAAAGAAAGCAAAATTAAGAGGAGAAAGAAATATAAATATAGAGTCT	599
Db	593	ATTATGTCAAAAGAAAGCAAAATTAAGAGAGAAAGAAATATAAATATAGAGTCT	652
Qy	600	TACTAGAAATTTGGCATCTAGAAATCTTGTCTCCAAAGGGAGAAATTCAGAAAAATATT	659
Db	653	TACTAGAAATTTGGCATCTAGAAATCTTGTCTCCAAAGGGAGAAATTCAGAAAAATATT	712
Qy	660	TACTGAGAAGTTAAGGAAGACAGTATTCCTGTCTCTGCTCTGTTGGCAGTATTAAT	719
Db	713	TACTGAGAAGTTAAGGAAGACAGTATTCCTGTCTCTGCTCTGTTGGCAGTATTAAT	772
Qy	720	CAACTTTACCCCTCGAGTATTTCCAAAGCTCTTCGTAATTCACAGTAGCAGAGAGGA	779
Db	773	CAACTTTACCCCTCGAGTATTTCCAAAGCTCTTCGTAATTCACAGTAGCAGAGAGGA	832
Qy	780	GGAGTGGCTACACAAACAGCTGAGGCACGAAGAGCAATGAATCTGACATAGCTGAAT	839
Db	833	GGAGTGGCTACACAAACAGCTGAGGCACGAAGAGCAATGAATCTGACATAGCTGAAT	892
Qy	840	TTGCGATTTAAAGAAAGAAAGAAACCCAGAAATGGTTGAAGGATAAAGAAACAAAT	899
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Qy	900	GTTTTCACAGCAAACTATTTGGCAGCTATCAATGCATATAATTTAGCCATAAGCTAAA	959
Db	953	GTTTTCACAGCAAACTATTTGGCAGCTATCAATGCATATAATTTAGCCATAAGCTAAA	1012
Qy	960	TAATAAGATGCCACTATTGTATTTGAACCGGGCTGCTTGCACACCTTAAACCTT	1019
Db	1013	TAATAAGATGCCACTATTGTATTTGAACCGGGCTGCTTGCACACCTTAAACCTT	1072
Qy	1020	ACACAGGCTATTGAAGATTCTTCTAAGGC	1049
Db	1073	ACACAGGCTATTGAAGATTCTTCTAAGGC	1102
RESULT 2			
LOCUS	CD107587	844 bp	mRNA linear EST 15-MAY-2003
DEFINITION	AGENCOURT 14021179 NIH_MGC_179 Homo sapiens cDNA clone		
	IMAGE:30368368 5', mRNA sequence.		
ACCESSION	CD107587		
VERSION	CD107587.1	GI:30760665	
KEYWORDS	EST.		
SOURCE	Homo sapiens (human)		
ORGANISM	Homo sapiens		
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.		
REFERENCE	1 (bases 1 to 844)		
AUTHORS	NIH-MGC http://mgc.nci.nih.gov/ .		
TITLE	National Institutes of Health, Mammalian Gene Collection (MGC)		
JOURNAL	Unpublished (1999)		
COMMENT	Contact: Robert Strausberg, Ph.D. Email: cgapbs-x@mail.nih.gov Tissue Procurement: Dr. Michael Brownstein CDNA Library Preparation: Invitrogen Corp CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL) DNA Sequencing by: Agencourt Bioscience Corporation Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov Plate: NDAM431 row: a column: 17 High quality sequence stop: 634.		
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		/lab_host="DHI08-Ton A (T1 and T5 phage resistances) "	
		/clone_lib="NIH_MGC_179"	
		/note="Organ: brain; Vector: pCMV-SPORT6.1; Site 1: EcoRV (destroyed); Site 2: NotI; Library is oligo-dT primed and directionally cloned (EcoRV site is destroyed upon cloning). Average insert size 1.1 kb. Library was constructed by (Invitrogen). Note: this is a NIH_MGC Library."	
ORIGIN			
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Qy	18	CGATTACAGCTGGCAGCAGACGAGACTGCGGTCTTCTGTCTCTGCCCCCTCAAAGGCGT	77
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Qy	78	GTGCGTCAGAGACACGACGCGTGTTCGACGGAAAACTATCTGAAGGTCAACTTTCCTCC	137
Db	61	GTGCGTCAGAGACACGACGCGTGTTCGACGGAAAACTATCTGAAGGTCAACTTTCCTCC	120
Qy	138	ATTTTATTTTGGAGCATTTCTTTATGCTCCCATAGACGATGAGAGCAAGCAAGAT	197
Db	121	ATTTTATTTTGGAGCATTTCTTTATGCTCCCATAGACGATGAGAGCAAGCAAGAT	180
Qy	198	TGGGAATGACACCACTTGTCTTACCTTGTATATAAAGAAAGCGCCATGTGGGAGACCT	257
Db	181	TGGGAATGACACCACTTGTCTTACCTTGTATATAAAGAAAGCGCCATGTGGGAGACCT	240
Qy	258	TTCTGTGACGCGGTGTTGACAAAGAGATGATCAAAAGAAATTAGAGAAAAATCTATTACA	317
Db			


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LOCUS             SILE0421p08f1 squirrel embryo library 1 Spermophilus lateralis
DEFINITION        cDNA clone 21p08 5', mRNA sequence.
ACCESSION         CO738958
VERSION           CO738958.1 GI:50826228
KEYWORDS          EST.
SOURCE            Spermophilus lateralis (golden-mantled ground squirrel)
ORGANISM          Spermophilus lateralis
                  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
                  Mammalia; Eutheria; Rodentia; Sciurognathi; Scuridae; Scurinae;
                  Spermophilus
                  1 (bases 1 to 793)
REFERENCE         Williams,D.R., Gracey,A.Y., Martin,S.L., Hughes,M.A., Li,W.;
                  Rogers,J. and Cossins,A.R.
                  Microarray analysis of transcriptional changes during hibernation
                  in the golden mantled ground squirrel, Spermophilus lateralis
                  Unpublished (2004)
JOURNAL           Contact: Andrew R. Cossins
COMMENT           Laboratory for Environmental Gene Regulation
                  University of Liverpool
                  School of Biological Sciences, The Biosciences Building, Crown
                  Street, Liverpool, United Kingdom, L69 7ZB
                  Tel: +44(0)151-795-4510
                  Fax: +44(0)151-795-4431
                  Email: cossins@liv.ac.uk
                  Vector has been trimmed from this EST.
                  Plate: 21 row: p column: 08
                  Seg primer: pflc T7 (5'-AATAGACTCACTATAGGG-3')
                  High quality sequence stop: 793.
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                    /lab_host="E.coli Electromax DH10B"
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                    GCATCC; Normalized and subtracted cDNA library prepared
                    from embryos"
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Best Local Similarity 90.3%; Pred. No. 7.5e-135;
Matches 714; Conservative 0; Mismatches 77; Indels 0; Gaps 0;
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Db 2 GAAGCAAAAGCTAGCAAGCGGGAAGATCAAAAATACGCACTAAGTGTCTCATGTAAG 61
Qy 406 ATTAGAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 465
Db 62 ATTAGAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 121
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Db 182 CAGAGAGAGAGAGAAATTATGTCAAAAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 241
Qy 586 AATATATAGAGCTCTTACTAGAAATTTGGCATCTAGAAATCTTGTCCCAAAAGGAGAAAT 645
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Qy 646 TCAGAAATAATATTTACTGAGAAAGTTAAAGGAGAGACAGTATTCTCTGCTCCTCGCTGTT 705
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Db 362 GGAATATTTAAATCAATTTTACCCCTCGAGTATTTCCCAACAGCTCTTCTGTAATCAAA 421
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Qy 946 GCCATAGAGCTAAATAATAAGATGCCACTATTGTTATTTGAACCGGGCTGCTTGGCAGCTA 1005
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Qy 1006 AAACATAAAACCTTTTACACAGAGCTTATTAAGATTTCTTAAAGCAGCTGGAATATTGATG 1065
Db 662 AAACATAAAACCTTTTACACAGAGCTTATTAAGATTTCTTAAAGCAGCTGGAATATTGATG 721
Qy 1066 CCACCTGTTACAGACATGCTTAATGCAAGATGAGGACACATGTACGACGTGGAGACAGCA 1125
Db 722 CCACATGTTGAGACAAATGCTTAATGCAAGATGAGGACACATGTACGACGTGGAGACAGCA 781
Qy 1126 TTCTGTCAACT 1136
Db 782 TTCTGTCAACT 792
RESULT 5
LOCUS             BG771796
DEFINITION        602720472P1 NIH_MGC_97 Homo sapiens cDNA clone IMAGE:4837505 5',
                  mRNA sequence.
ACCESSION         BG771796
VERSION           BG771796.1 GI:14082449
KEYWORDS          EST.
SOURCE            Homo sapiens (human)
ORGANISM          Homo sapiens
                  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
                  Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE         NIH-MGC http://mgi.nci.nih.gov/.
                  1 (bases 1 to 792)
AUTHORS           National Institutes of Health, Mammalian Gene Collection (MGC)
TITLE             Unpublished (1999)
JOURNAL           Contact: Robert Straubeberg, Ph.D.
COMMENT           Email: cgabbs-@mail.nih.gov
                  Tissue Procurement: Miklos Palkovits, M.D., Ph.D.
                  cDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki
                  Toshiyuki and Piero Carninci (RIKEN)
                  cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
                  DNA Sequencing by: Incyte Genomics, Inc.
                  Clone distribution: MGC clone distribution information can be
                  found through the I.M.A.G.E. Consortium/LLNL at:
                  http://image.llnl.gov
                  Plate: LLAM10769 Row: p column: 18
                  High quality sequence stop: 671.
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pBluescript KS+; Site 1: BamHI; Site 2: Sall-XhoI
 (gtcgag); Oligo-dT primed using primer
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 insert size 2.2 kb and normalized to ROT 5. This is a
 primary library enriched for full-length clones and
 constructed using the Cap-trapper method (Carninci, in
 preparation). Library constructed by M. Brownstein
 (NIMH/NHGRI, National Institutes of Health). Note: this is
 a NIH_MGC Library."

ORIGIN

Query Match 44.9%; Score 567.4; DB 4; Length 792;
 Best Local Similarity 91.5%; Pred. No. 5, 5e-113;
 Matches 671; Conservative 0; Mismatches 46; Indels 16; Gaps 6;

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 DB |||||
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 QY 181 AGCAGCAAGCAAGATTGGGAATGACACCATTTGTTTCTTACCTTGTATAAAAAAGAGCG 240
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 QY 241 GCATGTGGGAGACCTTTCTGTGAGGGTGTGTGACAAAGAGATGATGCAAGAAATTAGA 300
 DB |||||
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 QY 361 GCAGAGCGGAGAGATCAAAAAATACGCATTAAGTGTATGATGAAGATTGAAGAGAGAG 420
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 QY 637 AAGAGCTTACTAGAAAATTTGGCATCTAGAAATCTTGCTCCAGAGGGGAGAAATTC 696
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 QY 649 G--AAAATATATTTACTGAGAGTTAAGAGAGACAGTATTC--TGCTCCTCGTCTGT 704
 DB |||||
 QY 697 GCCCAATATATTTACTGAGAGTAAAGAGAGAAACAGTATTTCCCTGCTCCGTCGTGT 756
 DB |||||
 QY 705 TGGCAGTATTA 717
 DB |||||
 QY 757 TGGCAGACTTAA 769
 DB |||||

RESULT 6

BE564350
 LOCUS 601343161F1 NIH_MGC_53 Homo sapiens cDNA clone IMAGE:3685335 5',
 DEFINITION mRNA sequence.

ACCESSION

BE564350
 BE564350.1 GI:9808070

VERSION

EST.

KEYWORDS

SOURCE

ORGANISM

Homo sapiens (human)

REFERENCE

1 (bases 1 to 788)

NIH-MGC http://mgc.nci.nih.gov/.

National Institutes of Health, Mammalian Gene Collection (MGC)

Unpublished (1999)

Contact: Robert Strausberg, Ph.D.

Email: cgapbs-remail.nih.gov

Tissue Procurement: ATCC

cDNA Library Preparation: CLONTECH Laboratories, Inc.

cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)

DNA Sequencing by: Incyte Genomics, Inc.

Clone distribution: MGC clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at: image.llnl.gov

Plate: L1CM373 row: i column: 16

High quality sequence stop: 556.

Location/Qualifiers

1..788

source

/organism="Homo sapiens"

/mol_type="mRNA"

/db_xref="taxon:9606"

/clone="IMAGE:3685335"

/tissue type="carcinoma, cell line"

/lab host="DH10B (T1 phage-resistant)"

/clone lib="NIH MGC 53"

/note="Organ: bladder; Vector: pDNR-LIB (Clontech);

Site 1: SfiI (ggcgctcgcc); Site 2: SfiI

(ggccatattggcc); Double-stranded cDNA was prepared from

cell line RNA. 5' and 3' adaptors were used in cloning as

follows: 5' adaptor sequence: 5'-CACGCCATTATGCC-3' and

3' adaptor sequence:

5'-ATTCTAGAGCGGCGGCGGCACATG-dT(30)BN-3' (where B = A,

C, or G and N = A, C, G, or T). Average insert size 1.55

kb (range 0.9-4.0 kb). 15/15 colonies contained inserts

by PCR. This library was enriched for full-length clones

and was constructed by Clontech Laboratories (Palo Alto,

CA)."

ORIGIN

Query Match 43.8%; Score 553; DB 2; Length 788;

Best Local Similarity 94.6%; Pred. No. 7, 6e-110;

Matches 594; Conservative 0; Mismatches 31; Indels 3; Gaps 2;

QY 1 ATGCTCTTTCAGGTTAGCGATTACAGTGGCAGACGAGACATGCGGTCTTTCTGTCT 60

DB |||||

QY 23 ATGCTCTTTCAGGTTAGCGATTACAGTGGCAGACGAGACATGCGGTCTTTCTGTCT 82

DB |||||

QY 61 CTGCCCCCTCAAGGGCGTGTGCTGAGACACGAGCGGTCTGACCGAAAACTATCTG 120

DB |||||

QY 83 CTGCCCCCTCAAGGGCGTGTGCTGAGACACGAGCGGTCTGACCGAAAACTATCTG 142

DB |||||

QY 121 AAGGTCAACTTTCCTCCATTTTATTGAGGCATTTCTTTATGCTCCCATAGACGATGAG 180

DB |||||

QY 143 AAGGTCAACTTTCCTCCATTTTATTGAGGCATTTCTTTATGCTCCCATAGACGATGAG 202

DB |||||

QY 181 AGCAGCAAGCAAGATTGGGAATGACACCATTTGTCTTACCTTGTATATAAAGAGAGCG 240

DB |||||

QY 203 AGCAGCAAGCAAGATTGGGAATGACACCATTTGTCTTACCTTGTATATAAAGAGAGCG 262

DB |||||

QY 241 GCCATGTGGAGACCTTTCTGTGACGCGGTCTGACAAAGAGATGATGCAAGAAATTAGA 300

DB |||||

QY 263 GCCATGTGGAGACCTTTCTGTGACGCGGTCTGACAAAGAGATGATGCAAGAAATTAGA 322

DB |||||

QY 301 GAAAAATCTATTTTACAGCAAGAGAGAGCAAAAGAGCTTACAGAGCAAAAGCTGCA 360

DB |||||

QY 323 GAAAAATCTATTTTACAGCAAGAGAGAGCAAAAGAGCTTACAGAGCAAAAGCTGCA 382

DB |||||

QY 361 GCAAGCGGGAAGATCAAAAAATACGCATTAAGTGTATGATGAAGATTGAAGAGAGAG 420

DB |||||

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383 GCAAGCGGAGATCAAAATACGCCTAAGTCTCATGATGAGATTGACGCAAGAG 442
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421 AGG--AAAAAATAGAGATATGAAGAAATGAACCGGATAAAGCCACTAAAGCATTTGG 478
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443 AGGCAAAACAAATAGAGATATGAAGAAATGAACCGGATAAAGCCACTAAAGCATTTGG 502
    |||
479 AGGCTGGAGAGATATCAAGAAAGCTGAGGAGCAAAAAAATTCACAGAGAAAGAGA 538
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503 AGGCTGGAAAGAAATATCAAGAAAGCTGAGGAGCAAAAAAATTCAGAGAGAGAGA 562
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539 AATTATGTCAAAAAAGAAAGCAAAATTAAGAAAGCAAAAAAATTAATAATATAGAGTC 598
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563 ACTCTTGTG-CACAGAAAGCACTTCACGAGCGCGACACACCACCAATAAATATAGAGTC 621
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599 TTACTAGAAATTTGGCATCTAGAAATCT 626
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622 TTACTAGAAATGTGGCTTCTAGAAATCT 649
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RESULT 7
BP460416          683 bp      mRNA      linear      EST 31-DEC-2003
LOCUS             BP460416 full-length enriched swine cDNA library, adult ovary Sus
DEFINITION        BP460416 cDNA clone OVRM10147G01 5', mRNA sequence.
ACCESSION          BP460416
VERSION            BP460416.1 GI:40476478
KEYWORDS           EST.
SOURCE             Sus scrofa (pig)
ORGANISM           Sus scrofa

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REFERENCE
AUTHORS            Uenishi,H., Eguchi,T., Suzuki,K., Sawazaki,T., Toki,D., Shinkai,H.,
                   Okumura,N., Hamsima,N. and Awata,T.
TITLE              PEDE (Pig EST Data Explorer): construction of a database for ESTs
                   derived from porcine full-length cDNA libraries
JOURNAL            Nucleic Acids Res. 32 (1), D484-D488 (2004)
COMMENT            Contact: Hirohide Uenishi
                   Animal Genome Laboratory, Genome Research Department
                   National Institute of Agrobiological Sciences
                   2 Ikenodai, Tsukuba, Ibaraki 305-8602, Japan
                   Tel.: +81-29-838-8627
                   Fax: +81-29-838-8627
                   Email: huenishi@affrc.go.jp

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EST project with full-length enriched cDNA libraries carried out in
Animal Genome Research Program (Japan) by National Institute of
Agrobiological Sciences and STAFF-Institute
Single pass sequencing of clones derived from oligo-capped cDNA
library
Vector sequences were eliminated by RepeatMasker version 2002/07/13
and crossmatch version 0.990319
Low quality bases were trimmed based on the quality values.
FEATURES
source
Location/Qualifiers
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/organism="Sus scrofa"
/mol_type="mRNA"
/db_xref="taxon:9823"
/clone="OVRM10147G01"
/tissue_type="ovary"
/dev_stage="adult"
/clone_lib="full-length enriched swine cDNA library, adult
ovary"

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ORIGIN
Query Match          41.1%; Score 519; DB 5; Length 683;
Best Local Similarity 90.2%; Pred. No. 1.9e-102;
Matches 555; Conservative 0; Mismatches 60; Indels 0; Gaps 0;

539 AATTATGTCAAAAGAAAGCAAAATTAAGAAAGGAGGAAAAAATAAATAATATAGAGTC 598
|||||
1 AATTACATCAACAGGAAAGCAAAATTTGAAGAGAGAGAAAAAATTAAGCGGTAAAGGCC 60
|||||

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Qy 599 TTACTAGAAATTTGGCATCTAGAAATCTTCTGCCAAAGGGAGAGAAATTCAGAAATATAT 658
Db 61 TTACTAGAAATTTGAGCATCTAGATATCTTGACACAAATGGAGAAATTCAGAAATATAT 120
Qy 659 TTACTGAGAAAGTTAAAGGAGACAGTATCTCTGCTCTCTCGCTCTGTTGGCAGTATTTAAA 718
Db 121 TTTTGTGAGAAAGTTAAAGGAGACAGTATCTCTGCTCTCTCGCTCTGTTGGCAGTATTTAAA 180
Qy 719 TCACTTTTACCCCTCGAGTATTTCCACACAGCTCTTCTGATGATCACTCAAGTAGCAGCAAGG 778
Db 181 TCAACTTTTACCCCTCGAGTATTTCCCAACCGCCCTCCGGGAATCACAAGTAGCGGAAAG 240
Qy 779 AGAGTGGCTACACAAACAAAGCTGAGGCAGAGAGCAATGAATCTGACATAGCTGAAC 838
Db 241 AGAGTGGCTACACAAACAAAGCAGAGGCACGAGGGCAATGAATCTGAAATTCCTGAGT 300
Qy 839 TTTGCGAATTTAAAGAGAAAGAAACCCAGAAATGGTTGAAGGATAAAGGAAACAAAT 898
Db 301 TTAGTGATTTAAAGAGAGGAAAGAAACCCAGAAATGGTTGAAGGACACAAAGGAACAAAGT 360
Qy 899 TGTTTGCAACGGAAACTATTTGGCAGCTATCAATGCATATATTTAGCCATAGACTAA 958
Db 361 TGTTTGCAACGAGAAACTATTTGGCAGCTATTAATGCATACAACTTAGCCATAAGACTAA 420
Qy 959 ATAATAAGATGCCACTATTTGTTGAACCGGGCTGCTTGCCACTAAAACTAAAAAACC 1018
Db 421 ATAATAAGATTCACACTGTTGTTGATCGGGCTGCTTGCCACTAAAACTAAAAAACC 480
Qy 1019 TACAAGGCTATTGAAGATTTCTTAAGGACCTGGAATATTATGATGCCACCTGTTACAG 1078
Db 481 TACAAGGCTATTGAAGATTTCTTAAGGACCTGGAATATTATTAACACCACTGTTGAG 540
Qy 1079 ACATGCTTAATGCAAGATGAAGGCACATGTAGCAGCTGGAACAGCATTCCTGCAACTAG 1138
Db 541 ACAATGCTTAATGCAAGATGAAGGCACATGTAGCAGCTGGAACAGCATTCCTGCAACTAG 600
Qy 1139 AATTGTATGTAGAAG 1153
Db 601 AATTGTATGTAGAAG 615

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RESULT 8
AK005832          1190 bp      mRNA      linear      HTC 03-APR-2004
LOCUS             Mus musculus adult male testis cDNA, RIKEN full-length enriched
DEFINITION        library, clone:1700010i24 product:EKX1 homolog [Homo sapiens], full
                   insert sequence.
ACCESSION          AK005832
VERSION            AK005832.1 GI:12838612
KEYWORDS           HTC; CAP trapper.
SOURCE             Mus musculus (house mouse)
ORGANISM           Mus musculus

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REFERENCE
AUTHORS            Carninci,P. and Hayashizaki,Y.
TITLE              High-efficiency full-length cDNA cloning
JOURNAL            Meth. Enzymol. 303, 19-44 (1999)
MEDLINE            99279253
PUBMED             10349636

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REFERENCE
AUTHORS            Carninci,P., Shibata,Y., Hayatsu,N., Sugahara,Y., Shibata,K.,
                   Itoh,M., Kono,H., Okazaki,Y., Muramatsu,M. and Hayashizaki,Y.
TITLE              Normalization and subtraction of cap-trapper-selected cDNAs to
                   prepare full-length cDNA libraries for rapid discovery of new genes
JOURNAL            Genome Res. 10 (10), 1617-1630 (2000)
MEDLINE            20499374
PUBMED             11042159
AUTHORS            Shibata,K., Itoh,M., Aizawa,K., Nagaoaka,S., Sasaki,N., Carninci,P.,
                   Konno,H., Akiyama,J., Nishi,K., Kitsuai,T., Tashiro,H., Itoh,M.,
                   Sumi,N., Ishii,Y., Nakamura,S., Hazama,M., Nishine,T., Harada,A.,
                   Yamamoto,R., Matsumoto,H., Sakaguchi,S., Ikegami,T., Kashiwagi,K.,

```

Fujiwaka, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E., Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
 RIKEN integrated sequence analysis (RISA) system--384-format
 sequencing pipeline with 384 multicapillary sequencer
 Genome Res. 10 (11), 1757-1771 (2000)
 20530913
 11076861

REFERENCE

4 The RIKEN Genome Exploration Research Group Phase II Team and the FANTOM Consortium.

Functional annotation of a full-length mouse cDNA collection

JOURNAL

Nature 409, 685-690 (2001)

REFERENCE

5 The FANTOM Consortium and the RIKEN Genome Exploration Research

Group Phase I & II Team.

Analysis of the mouse transcriptome based on functional annotation

JOURNAL

of 60,770 full-length cDNAs

REFERENCE

Nature 420, 563-573 (2002)

AUTHORS

6 (bases 1 to 1190)
 Adachi, J., Aizawa, K., Akahira, S., Akimura, T., Arai, A., Aono, H., Arakawa, T., Bono, H., Carninci, P., Fukuda, S., Fukunishi, Y., Furuno, M., Hanagaki, T., Hara, A., Hayatsu, N., Hiramoto, K., Hiraoka, T., Hori, F., Imotani, K., Ishii, Y., Itoh, M., Izawa, M., Kasukawa, T., Kato, H., Kawai, J., Kojima, Y., Konno, H., Kouda, M., Koya, S., Kurihara, C., Matsuyama, T., Miyazaki, A., Nishi, K., Nomura, K., Numazaki, R., Ohno, M., Okazaki, Y., Okido, T., Owa, C., Saito, H., Saito, R., Sakai, C., Sakai, K., Sano, H., Sasaki, D., Shibata, K., Shibata, Y., Shinagawa, A., Shiraki, T., Sogabe, Y., Suzuki, H., Tagami, M., Tagawa, A., Takahashi, F., Tanaka, T., Tejima, Y., Toya, T., Yamamura, T., Yasunishi, A., Yoshida, K., Yoshino, M., Muramatsu, M. and Hayashizaki, Y.

TITLE

Submitted (10-JUL-2000) Yoshihide Hayashizaki, The Institute of Physical and Chemical Research (RIKEN), Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), RIKEN Yokohama Institute, 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan (E-mail: genome-res@gsc.riken.jp, URL: <http://genome.gsc.riken.jp/>, Tel: 81-45-503-9222, Fax: 81-45-503-9216)

COMMENT

Please visit our web site (<http://genome.gsc.riken.jp/>) for further details.

cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. First strand cDNA was primed with a primer [5' GAGAGAGAGAGATCCAGAGCTCTTTTCTTTTNN 3'], cDNA was prepared by using trehalose thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. Second strand cDNA was prepared with the primer adapter of sequence [5' GAGAGAGAGAGCGCCCAATTAACTTCGAGTTAAATAATCCGCCCC 3']. cDNA was cleaved with XhoI and SstI. Cloning sites, 5' end: XhoI, 3' end: SstI. Host: SOUR.

FEATURES

source

Location/Qualifiers
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 /tissue_type="testis"
 /clone_lib="RIKEN full-length enriched mouse cDNA library"
 /dev_stage="adult"
 351..803
 /note="unnamed protein product; EK1 homolog [Homo sapiens] (SPTR|AAL73230, evidence: FASTY, 89.1%id, 41.4%length, match=522)
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/db_xref="GI:26348135"
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 1169..1174
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ORIGIN

Query Match 40.7%; Score 513.8; DB 3; Length 1190;
 Best Local Similarity 79.5%; Pred. No. 2.8e-101;
 Matches 634; Conservative 0; Mismatches 157; Indels 6; Gaps 2;
 QY 464 CCATTAAGCATTGGAGCCTGGAAAGAAATATCAAGAAAGAGCTGAGAGCAAAAAA 523
 Db 111
 4 CAATAGCGAATTAGAGCGTGGAAAGAAATGTCAAAAGAAAGCTGACGGAACAAAAGAG 63
 QY 524 TTCAGAGAGAGAGAAATTTATGTCAAAAGAAAGAAAGAAATTAAGAGAGAGAAAA 583
 Db 111
 64 TCCAGAGAGAGAGAA---ACCGCTCGAGGAGAAAGCAAGCTGAGAG---ACCAAAGCTC 117
 QY 584 TAAATATAAGAGTCTTACTAGAAATTTGGCATCTAGAAATCTTGCTCCAAAAGGGAGAA 643
 Db 111
 118 TAAACCTCGGGTTTGGCCCGGAGGCCCCACCCACTCGCTCCCAACAGAGGGAGGA 177
 QY 644 ATTCAAGAAATATATTTATCTGAGAGTGTAAAGGAGAGACAGTATTTCTGCTCTG 703
 Db 111
 178 ATTGGGAAACATATTTCTCTGAGAGTGTAAAGGAGAGACAGAGTCCCTCGCTCG 237
 QY 704 TTGGCAGTATTAATAATCACTTTACCTTCAGTATTTCCCAACAGCTCTTCTGTGATCAC 763
 Db 111
 238 CTGGCAGTATTCAAATCAGCTTTACCTTCGAGTGTTCCTCAACAGCACTTCGGGAATCCC 297
 QY 764 AAGTAGCAGAGAGAGAGAGTGGCTACACAAAGCTGAGGAGAGAGCAATGAATA 823
 Db 111
 298 AAGTCGAGAGAGAGAGAGTGGCTGCATTAACAGAGAGAGAGAGAGAGAGAGAGCA 357
 QY 824 CTGACATAGCTGAACCTTTGCGATTTAAAGAGAGAGAGAGAGAGAGAGAGAGAGAG 883
 Db 111
 358 CTGACCTTCTCTGAGTCTTTTGACTTTAAAGAGAGAGAGAGAGAGAGAGAGAGAG 417
 QY 884 ATAAAG 943
 Db 111
 418 ACAAGGGAG 477
 QY 944 TAGCCATAAG 1003
 Db 111
 478 TAGCCATAG 537
 QY 1004 TAAAG 1063
 Db 111
 538 TCAATTAAG 597
 QY 1064 TGCCACCTGTTACAG 1123
 Db 111
 598 CACCACCTGTTGCTGCAATGCAATGCAATGCAATGCAATGCAATGCAATGCAATGCA 657
 QY 1124 CATTCGTCTGAG 1183
 Db 111
 658 CGTTCTGTCAACTAGAGATTTGATGTTGAAGGCTTGCAGAGATTTATGAAGCTGCAC 717
 QY 1184 TTGATTCATCAACAAATTTGATGATGATGATGATGATGATGATGATGATGATGATG 1243
 Db 111
 718 TTGACCCAGCCACACAGTGTGACAGACGATGACAGAGAGATTCGGNATATATTCAG 777
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 Db 111
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RESULT 9
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LOCUS BQ217312 1131 bp mRNA linear EST 02-MAY-2002
 DEFINITION AGENCOURT 7558431 NIH_MGC_72 Homo sapiens cDNA clone IMAGE:6047052
 5', mRNA sequence.
 ACCESSION BQ217312
 VERSION BQ217312.1 GI:20398712
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 NIH-MGC <http://mgc.nci.nih.gov/>.
 REFERENCE 1 (bases 1 to 1131)
 AUTHORS National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL Unpublished (1999)
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgabs-remail.nih.gov
 Tissue Procurement: ATCC/DCFT/DTF
 CDNA Library Preparation: Life Technologies, Inc.
 CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Agencourt Bioscience Corporation
 Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
 Plate: L1AM13293 row: n column: 13
 High quality sequence stop: 388.
 Location/Qualifiers
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 /clone="IMAGE:6047052"
 /tissue_type="melanotic melanoma"
 /lab_host="DH10B (phage-resistant)"
 /clone_lib="NIH MGC 72"
 /note="Organ: skin; Vector: pCMV-SPORT6; Site 1: NotI; Site 2: SalI; Cloned unidirectionally. Primer: Oligo dT. Average insert size 2 kb. Library constructed by Life Technologies."
 FEATURES
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 Query Match 40.4%; Score 510.4; DB 5; Length 1131;
 Best Local Similarity 95.9%; Pred. No. 1.5e-100;
 Matches 535; Conservative 0; Mismatches 21; Indels 2; Gaps 1;
 Qy 42 GACTGGGCTCTTCTGCTCTGCTCCCTCAAGGGGTGCTGCTGAGACGACGACGACGCTGT 101
 Db 1 GACTGGGCTCTTCTGCTCTGCTCCCTCAAGGGGTGCTGCTGAGACGACGACGACGCTGT 60
 Qy 102 CTGACGGAAACTATCTGAGGTCACCTTCTCCCTTTTATTTGAGGCTTCTTTA 161
 Db 61 CTGACGGAAACTATCTGAGGTCACCTTCTCCCTTTTATTTGAGGCTTCTTTA 120
 Qy 162 TGCTCCCATAGACGATGAGCAGCAAGCAAGATGGGAATGACACCATTTGCTTCAC 221
 Db 121 TGCTCCCATAGCAGTATGAGCAGCAAGCAAGATGGGAATGACACCATTTGCTTCAC 180
 Qy 222 CTTGTATATAAAGAACGGCCATGTGGGACCCCTTCTGTGCGGTGTGCAAGA 281
 Db 181 CTTGTATATAAAGAACGGCCATGTGGGACCCCTTCTGTGCGGTGTGCAAGA 240
 Qy 282 GATGATGCAAGAACTAGAGAAATCTATTTTACAAGCACAAGAGAGCAAAAGAGC 341
 Db 241 GATGATGCAAGAACTAGAGAAATCTATTTTACAAGCACAAGAGAGCAAAAGAGC 300
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 Db 301 TACAGACCAAAAGCTGACGAAACGGGAAGATCAAAATACGCACTTAAGTGCATCAT 360
 Qy 402 GAAGATTGAAGAAAGAGAGGAAATAATAGAGATATGAAGAAATGAACGGATAAA 461
 Db 361 GAAGATTGAAGAAAGAGAGGAAATAATAGAGATATGAAGAAATGAACGGATAAA 420
 Qy 462 AGCCACTAAGCATTTGGAAGCTTGAAAGAAATATCAAGAAAGCTGAGGAGC--AAAAA 519

Db 421 AGCCACTAAGCATTTGGAAGCTTGACAGAGATATCAAGAAAGAGCTGAGGAGCAAAAAA 480
 Qy 520 AAAATTCAGAGAGAGAGAAATTTATGTCAAAAGAGAGCAAAATTAAGAGAGGAGAAA 579
 Db 481 AAAATTCAGAGAGAGAGAAATTTATGTCAAAAGAGAGAAACCCCAATTAAGAGAAATGAAGGA 540
 Qy 580 AAAATTAATAATTAAGACT 597
 Db 541 AAAAAAATAATAATAGT 558
 RESULT 10
 LOCUS BY705981
 DEFINITION BY705981 RIKEN full-length enriched, adult male testis Mus musculus cDNA clone 1700010124 5', mRNA sequence.
 ACCESSION BY705981
 VERSION BY705981.1 GI:27117128
 KEYWORDS EST.
 SOURCE Mus musculus (house mouse)
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 917)
 Okazaki, Y., Furuno, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S., Nikaido, I., Osato, N., Saito, R., Suzuki, H., Yamanaka, I., Kiyosawa, H., Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A., Schonbach, C., Gojobori, T., Baldarelli, R., Hill, D. P., Bult, C., Hume, D. A., Quackenbush, J., Schriml, L. M., Kanapin, A., Matsuda, H., Batalov, S., Beisel, K. W., Blake, J. A., Bradt, D., Bruscia, V., Chothia, C., Corbani, L. E., Cousins, S., Dalla, E., Dragani, T. A., Fletcher, C. F., Forrest, A., Frazer, K. S., Gaasterland, T., Gariboldi, M., Glissi, C., Godzik, A., Gough, J., Grimmond, S., Gustincich, S., Hirokawa, N., Jackson, I. J., Jarvis, E. D., Kanai, A., Kawaji, H., Kawasawa, Y., Kedzierzki, R. M., King, B. L., Konagaya, A., Kurochkin, I. V., Lee, Y., Lenhard, B., Lyons, P. A., Maglott, D. R., Maltais, L., Marchionni, L., McKenzie, L., Miki, H., Nagashima, T., Numata, K., Okido, T., Pavan, W. J., Pertea, G., Pesole, G., Petrovsky, N., Pillai, R., Pontius, J. U., Qi, D., Ramachandran, S., Ravasi, T., Reed, J. C., Reid, J., Ring, B. Z., Ringwald, M., Sandelin, A., Schneider, C., Semple, C. A., Setou, M., Shimada, K., Sultana, R., Takenaka, Y., Taylor, M. S., Teasdale, R. D., Tomita, M., Varado, R., Wagner, L., Wahlestedt, C., Wang, Y., Watanabe, Y., Wells, C., Wilming, L. G., Wynshaw-Boris, A., Yanagisawa, M., Yang, I., Yang, D., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A., Carninci, P., Hayatsu, N., Hirozane-Kishikawa, T., Konno, H., Nakamura, M., Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Aizawa, K., Arakawa, T., Fukuda, S., Hara, A., Hashizume, W., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata, K., Shinagawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Lander, E. S., Rogers, J., Birney, E. and Hayashizaki, Y.
 Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs
 Nature 420, 563-573 (2002)
 22354683
 1246851
 Contact: Yoshihide Hayashizaki
 Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute
 The Institute of Physical and Chemical Research (RIKEN)
 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
 Tel: 81-45-503-9222
 Fax: 81-45-503-9216
 Email: genome-res@gsc.riken.jp, [URL:http://genome.gsc.riken.jp/](http://genome.gsc.riken.jp/)
 Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda, S., Hashizume, W., Hayashida, K., Hirozane, T., Hori, F., Imotani, K., Ishii, Y., Itoh, M., Kagawa, I., Kawai, J., Kojima, Y., Kondo, S., Komori, H., Koya, S., Miyazaki, A., Murata, M., Nakamura, M., Nomura, K., Numazaki, R., Ohno, M., Ohsato, N., Saito, R., Sakazume, N., Sano, H., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami, M., Takeda, Y., Waki, K., Watahiki, A., Muramatsu, M. and Hayashizaki, Y.
 Direct Submission


```

QY      866 ACCC 869
DB      514 AGCC 517

RESULT 13
LOCUS   CK603982
DEFINITION AGENCOURT_17890681 NIH_MGC_238 Rattus norvegicus cDNA clone
IMAGE: 7134374 5', mRNA sequence.
ACCESSION CK603982
VERSION   CK603982.1 GI:41117319
KEYWORDS EST.
SOURCE   Rattus norvegicus (Norway rat)
ORGANISM Rattus norvegicus
          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
          Rattus.
REFERENCE 1 (bases 1 to 796)
AUTHORS  NIH-MGC http://mgc.nci.nih.gov/.
TITLE    National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL  Unpublished (1999)
COMMENT  Contact: Daniela S. Gerhard, Ph.D.
          Office of Cancer Genomics
          National Cancer Institute / NIH
          Bldg. 31 Rm10A07 Bethesda, MD 20892
          Email: cgapbs-r@mail.nih.gov
          Tissue Procurement: Howard Jacobs
          cDNA Library Preparation: Express Genomics
          cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
          DNA Sequencing by: Agencourt Bioscience Corporation
          Clone distribution: MGC clone distribution information can be
          found through the I.M.A.G.E. Consortium/LLNL at:
          http://image.llnl.gov
          Plate: LLAM15039 row: o column: 12
          High quality sequence stop: 622.
          Location/Qualifiers
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              /mol_type="mRNA"
              /db_xref="taxon:10116"
              /clone="IMAGE:7134374"
              /issue_type="testis, pooled"
              /lab_host="DH10B Tona"
              /clone_lib="NIH MGC 238"
              /note="Organ: testis; Vector: pExpress-1; Site 1: EcoRV;
              Site 2: NotI; RNA obtained from testis tissue of 8 wk old
              animal. Tissues were snap-frozen and kept at -80C before
              RNA extraction and purification (TRI-reagent method). cDNA
              was primed using oligo-dT primer:
              5'-pGACTAGTCTAGATCGGCGGCGGCC(T)25-3' and cloned into
              the EcoRV/NotI sites of pExpress-1. Size-selection >1.4kb
              resulted in an average insert size of 1.9 kb. This primary
              library is normalized (non-normalized primary library is
              NIH_MGC_237) and was constructed by Express Genomics
              (Frederick, MD)"

ORIGIN
Query Match 35.6%; Score 449; DB 7; Length 796;
Best Local Similarity 76.9%; Pred. No. 3.4e-87;
Matches 574; Conservative 0; Mismatches 166; Indels 6; Gaps 2;

QY      1 ATGCGCTCTTCAGGTAGCATATACAGTGGCAGACGAGACGACCTGCGGTCTTTCTGTCT 60
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DB      4 ATGCGGTGCGAGTAGCGAGCTCAGTGGCAGACGACACCGCGCGCACTCTTCTGTCTG 63
QY      61 CTGCGCCCTCAAGGCGGTGCGTTCAGACACGAGCGTTCCTGCAACGGAACAACTATCTG 120
DB      |||||
DB      64 CTGCGCTACGGGGCGTCTGCGCGCATGCTGATGTTCTGTGGGGAAGTTACCTG 123
QY      121 AGGTCAACTTTCCTCCATTTTATTGAGGCATTTCTTATGCTCCCATAGACATGAG 180
DB      |||||
DB      124 AAGGTTAACTTTCCTCCATTTTATTGAGGTTTCTCTATGCTCCCATAGATGCGG 183

181 AGCAGCAAGCAAGATTGGGAATGACACCAATTGTCTTCACTTGTATAAAAAAGAGCG 240
184 AAGAGCAAGCAAGATTGGGAATGACACCAATTGTCTTCACTTGTATAAAAAAGAGCG 243
241 GCCATGTGGGAGACCCCTTTCTGTGACGGGTGTGACAAAGAGATGATGCARAGAATTAGA 300
244 GTTCTGTGGGAGAGCCCTTTCTATGCCAGCGTTGTATAAGAGATGATGCAGAGAATAAGA 303
301 GAAAAATCTATTTTACAAGCAACAAGAGAGAGCAAAAGAGCTTACAGAAAGCTGCA 360
304 GAAAAATCTATCTTGTCAAGCAGAGAGAAAGCAAAAGAGGCTTACAGAGCGAAGCTGCT 363
361 GCAAGCGGGAAGATCAAAAAATACCCACTAAGTGTCTATGATGAAGATTGAAGAAGAAG 420
364 GCCAAGCCGAGAAGATCAGAGATACGCCCTTAGCCGAGATGATGAAGATTGAAGAAGAAG 423
421 AGGAAAAAATAGAGATATGAAGAAATGAACCGATGAAGCAACCACTAAAGCAATTTGAA 480
424 AGGAAAAAATAGAGATATGAAGAAATGAACCGAATAATGAACGAAAAAGCAACCAAGAA 483
481 GCCTGGAAGAAATATCAAGAAAGAAAGCTGAGGAGCAAAAAAATTCAGAGAGAGAGAAA 540
484 CGGTGGAAGAAATGCCAAAGAAAGCTGACGGAACAAAACGAGTCCAGAGGAGAGAGAA 542
541 TTATGTCAAAAAGAAAGCAAAATTAAGAAAGAAAGAAAAAATAAATAATATAAGAGTCTT 600
543 --ACCGCTACAGGGGAAAGCAAGCT---GAAGAGAGGGGAGCACTAAAAACCTCAGAGTTTG 597
601 ACTAGAAATTTGGCATCTAGAAATCTTCTCCAAAGGAGGAGAAATTCAGAAATATATTT 660
598 CCCCAGAGGGCCCGCCCACTCGCCCTCCCAAGAGGAGGAAATTTGGGAAAAACATATTC 657
661 ACTGAGAAAGTTAAAGGAAGACAGTATTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 720
658 TCTGAGAGTTTAAAGGAAGACAGGGTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCAG 717
721 AACTTTACCCCTCGAGTATTCACAC 746
718 AGCTTTNACCCTCGAGTGTCCAC 743

CB297042 468 bp mRNA linear EST 28-FEB-2003
12B22058_rev_1_F09_r_075.ab1 Chimpanzee brain library Koo's Pan
troglodytes cDNA clone 12B22058_rev_1_F09_r_075.ab1 5', mRNA
sequence.
ACCESSION CB297042
VERSION   CB297042.1 GI:28622472
KEYWORDS EST.
SOURCE   Pan troglodytes (chimpanzee)
ORGANISM Pan troglodytes
          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
          Hellmann, I., Zollner, S., Enhard, W., Ebersberger, I., Nickel, B. and
          Paabo, S.
          Selection on human genes as revealed by comparisons to chimpanzee
          cDNA
          Genome Res. (2003) In press
          Contact: Paabo S
          Evolutionary Genetics
          Max-Planck-Institute for Evolutionary Anthropology
          Deutscher Platz 6, 04103 Leipzig, Germany
          Tel: +49-(0)-341-3550 500
          Fax: +49-(0)-341-3550 555
          Email: paabo@eva.mpg.de
          Seq primer: M13 reverse.
          Location/Qualifiers
            1..468
              /organism="Pan troglodytes"
              /mol_type="mRNA"

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Job time : 4610 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: July 8, 2005, 01:40:53 ; Search time 245 Seconds
(without alignments)
8435.173 Million cell updates/sec

Title: US-10-681-199-1
Perfect score: 1263
Sequence: 1 atgcctcttcaggttagcga.....gaacagaaactaaattcttaa 1263

Scoring table: IDENTITY NUC
Gapop 10.0 , Gapext 1.0

Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents NA.*
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2: /cgn2_6/ptodata/1/ina/5B_COMB.seq.*
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6: /cgn2_6/ptodata/1/ina/backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description	
1	432.6	34.3	489	4	US-09-621-976-1234	
2	178.8	14.2	313	4	Sequence 1234, Ap	
3	162.8	12.9	164	4	Sequence 30874, A	
C	4	115.8	9.2	7218	1	Sequence 34573, A
	5	90.8	7.2	612	4	Sequence 14, Appl
	6	77.4	6.1	1039	4	Sequence 1357, Ap
	7	77.2	6.1	614	4	Sequence 1280, Ap
8	73.4	5.8	258775	4	Sequence 1318, Ap	
9	72.6	5.7	705	4	Sequence 16435, A	
10	72.6	5.7	705	4	Sequence 5061, Ap	
11	72.4	5.7	240	1	Sequence 20343, A	
C	12	72	5.7	396	4	Sequence 6, Appl
	13	72	5.7	396	4	Sequence 53, Appl
C	14	72	5.7	396	4	Sequence 53, Appl
C	15	72	5.7	396	4	Sequence 53, Appl
16	71.2	5.6	1696	4	Sequence 53, Appl	
17	71.2	5.6	1827	4	Sequence 1, Appl	
18	71.2	5.6	1827	4	Sequence 1308, Ap	
19	70.2	5.6	851	4	Sequence 16590, A	
C	20	70.2	5.6	851	4	Sequence 8286, Ap
C	21	69.8	5.5	161124	4	Sequence 23568, A
22	69.4	5.5	2447	2	Sequence 11760, A	
C	23	67.4	5.3	25431	4	Sequence 14, Appl
	24	67.2	5.3	601	4	Sequence 13234, A
	25	67.2	5.3	601	4	Sequence 184917, A
	26	66.6	5.3	194937	4	Sequence 185072, A
27	66.6	5.3	194937	4	Sequence 17032, A	
28	66.6	5.3	194937	4	Sequence 17033, A	

28	66.2	5.2	696	3	US-09-461-697-193	Sequence 193, App	
29	66.2	5.2	699	3	US-09-461-697-191	Sequence 191, App	
30	66.2	5.2	717	3	US-09-461-697-189	Sequence 189, App	
31	66.2	5.2	774	3	US-09-461-697-187	Sequence 187, App	
32	66.2	5.2	819	3	US-09-461-697-185	Sequence 185, App	
33	66.2	5.2	1669	3	US-09-461-697-184	Sequence 184, App	
34	66.2	5.2	19124	2	US-08-487-826B-13	Sequence 13, Appl	
35	65.4	5.2	36731	4	US-09-949-016-13770	Sequence 13770, A	
36	64.6	5.1	2223	1	US-08-257-073-4	Sequence 4, Appli	
37	63.6	5.0	980	3	US-09-171-209-8	Sequence 8, Appli	
C	38	63.6	5.0	55841	4	US-09-949-016-16602	Sequence 16602, A
39	63.6	5.0	74790	4	US-09-949-016-15321	Sequence 15321, A	
40	63.4	5.0	5433	3	US-08-929-329-1	Sequence 1, Appli	
C	41	63.4	5.0	12703	4	US-09-949-016-16685	Sequence 16685, A
42	62.6	5.0	237510	4	US-09-949-016-14273	Sequence 14273, A	
C	43	62.4	4.9	54033	4	US-09-949-016-12091	Sequence 12091, A
C	44	62.4	4.9	54033	4	US-09-949-016-14325	Sequence 14325, A
45	62	4.9	1298	3	US-08-948-705-3	Sequence 3, Appli	

ALIGNMENTS

RESULT 1
US-09-621-976-1234
; Sequence 1234, Application US/09621976
; Patent No. 6639063
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Jobert, S.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: ESTs and Encoded Human Proteins.
; FILE REFERENCE: GENSET.054PR2
; CURRENT APPLICATION NUMBER: US/09/621.976
; CURRENT FILING DATE: 2000-07-21
; NUMBER OF SEQ ID NOS: 19335
; SOFTWARE: Patent.pm
; SEQ ID NO 1234
; LENGTH: 489
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 57..488
US-09-621-976-1234

Query Match		34.3%	Score 432.6;	DB 4;	Length 489;
Best Local Similarity		99.8%	Pred. No. 4.6e-97;		
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Qy	1	ATGCTCTTTCAGGTTAGCGATTACAGCTGGCAGACGAGAGACTGGCGTCTTTCTGTCT	60		
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Qy	61	CTGCCCTCAAGCGGTGGCTCAGACACGAGCGTTCTCGCGGAAACTATCTG	120		
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Qy	121	AAGTCAACTTCTCCATTTTATTTAGGCACTTTCTTTATGTCCCATAGACGATCAG	180		
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Qy	181	AGCAGCAAGCAAGATTGGGAATGACACCACTTCTTACCTTTGTATATAAAGAACGC	240		
Db	237	AGCAGCAAGCAAGATTGGGAATGACACCACTTCTTACCTTTGTATATAAAGAACGC	296		
Qy	241	GCCATGTGGGAGACCCCTTTCTGTACCGGTGTTGACAAAGAGATGATCAAGAAATAGA	300		
Db	297	GCCATGTGGGAGACCCCTTTCTGTACCGGTGTTGACAAAGAGATGATCAAGAAATAGA	356		
Qy	301	GAAAAATCTATTTTACAGCACAAGAGAGCAAGAAAGCTACAGAGCAAGAAAGCTGCA	360		
Db	357	GAAAAATCTATTTTACAGCACAAGAGAGCAAGAAAGCTACAGAGCAAGAAAGCTGCA	416		

Qy 361 GCAAGCGGAGATCAAAATACGCACTAAGTCTCATGATGAAGATTGAAGAAGAG 420
Db 417 GCAAGCGGAGATCAAAATACGCACTAAGTCTCATGATGAAGATTGAAGAAGAG 476
Qy 421 AGGAAAAAATAG 433
Db 477 AGGAAAAAATAG 489

RESULT 2
US-09-513-999C-30874
; Sequence 30874, Application US/09513999C
; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Duclert, A.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
; Patent No. 6783961
; FILE REFERENCE: 59.US2.REG
; CURRENT APPLICATION NUMBER: US/09/513.999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/122,487
; PRIOR FILING DATE: 1999-02-26
; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
; SEQ ID NO 30874
; LENGTH: 313
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 82
; OTHER INFORMATION: r=a or g
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 84
; OTHER INFORMATION: w=a or t
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 85
; OTHER INFORMATION: w=a or t
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 89
; OTHER INFORMATION: v=a or c or g
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 92
; OTHER INFORMATION: w=a or t
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 250
; OTHER INFORMATION: n=a, g, c or t
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 283
; OTHER INFORMATION: s=g or c
US-09-513-999C-30874

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Best Local Similarity 94.8%; Pred. No. 1.5e-34;
Matches 201; Conservative 5; Mismatches 3; Indels 3; Gaps 2;
Qy 844 GATTAAAGAGAGAGAAAGAACCCAGATGGTTGAAGGATAAGGAAACAAATTGTTT 903
Db 1 GATTAAAGAGAGAGAGAAAGAACCCAGATGGTTGAAGGATAAGGAAACAAATTGTTT 60
Qy 904 GCAACGGAAACTATTGGCGAGCTA--TCAATGCATAT-AATTTAGCCATAAGCTAAAT 960
Db 61 GCAACGGAAACTATTGGCGARCWATCAWAGCATATAAATTTAGCCATAAGCTAAAT 120

Qy 961 AATAAGATGCCACTATTGTATTGAACCGGCTGCTTGCACCTAAACCTAAAAACTTA 1020
Db 121 AATAAGATGCCACTATTGTATTGAACCGGCTGCTTGCACCTAAACCTAAAAACTTA 180
Qy 1021 CACAAGCTATTGAAGATTCTTCTTAAGGCACT 1052
Db 181 CACAAGCTATTGAAGATTCTTCTTAAGGAATT 212

RESULT 3
US-09-513-999C-34573
; Sequence 34573, Application US/09513999C
; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Duclert, A.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
; Patent No. 6783961
; FILE REFERENCE: 59.US2.REG
; CURRENT APPLICATION NUMBER: US/09/513.999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/122,487
; PRIOR FILING DATE: 1999-02-26
; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
; SEQ ID NO 34573
; LENGTH: 164
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 127
; OTHER INFORMATION: y=c or t
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 128
; OTHER INFORMATION: h=a or c or t
US-09-513-999C-34573

Query Match 12.9%; Score 162.8; DB 4; Length 164;
Best Local Similarity 98.8%; Pred. No. 9.9e-31;
Matches 162; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Qy 5 CTCTCAGGTTAGCATTTACAGCTGGCAGCAGACGAGACTGCGGTCTTTCTGTCCTGC 64
Db 1 CTCTCAGGTTAGCATTTACAGCTGGCAGCAGACGAGACTGCGGTCTTTCTGTCCTGC 60
Qy 65 CCCTCAAAGGCGTGTGCTCAGAGACACGACGCTGTTCTGCACGGAAACCTATCTGAAGG 124
Db 61 CCCTCAAAGGCGTGTGCTCAGAGACACGACGCTGTTCTGCACGGAAACCTATCTGAAGG 120
Qy 125 TCAACTTCTCCATTTTATTTGAGGCATTTCTTTATGCTCCC 168
Db 121 TCAACTVHCCTCCATTTTATTTGAGGCATTTCTTTATGCTCCC 164

RESULT 4
US-08-232-463-14/c
; Sequence 14, Application US/08232463
; Patent No. 5670367
; GENERAL INFORMATION:
; APPLICANT: DORNER, F.
; APPLICANT: SCHEIFLINGER, F.
; APPLICANT: FALKNER, F. G.
; TITLE OF INVENTION: RECOMBINANT FOWLPOX VIRUS
; NUMBER OF SEQUENCES: 52
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Foley & Lardner
; STREET: 1800 Diagonal Road, Suite 500
; CITY: Alexandria
; STATE: VA
; COUNTRY: USA

Qy 574 AGAAAAATAATAATATAAG 596
Db 122787 AAGAAAGAAAGAAAGAAAG 122809

RESULT 9
US-09-270-767-5061
; Sequence 5061, Application US/09270767
; Patent No. 6703491
; GENERAL INFORMATION:
; APPLICANT: Homburger et al.
; TITLE OF INVENTION: Nucleic acids and proteins of *Drosophila melanogaster*
; FILE REFERENCE: File Reference: 7326-094
; CURRENT APPLICATION NUMBER: US/09/270,767
; CURRENT FILING DATE: 1999-03-17
; NUMBER OF SEQ ID NOS: 62517
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 5061
; LENGTH: 705
; TYPE: DNA
; ORGANISM: *Drosophila melanogaster*
US-09-270-767-5061

Query Match 5.7%; Score 72.6; DB 4; Length 705;
Best Local Similarity 51.8%; Pred. No. 3.1e-08;
Matches 216; Conservative 0; Mismatches 194; Indels 7; Gaps 2;

Qy 264 GACGGGTGTTCCACAAAGAGATGATGCTCAAAGAATTAGAGAAAAATCTATTTTACAAGCACA 323
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Qy 324 AGAGAGAGCAAAAGAAAGCTACAGAAAGCAAAAAGCTGCAGCAAGCGGGGAAGATCAAAAATA 383
Db 231 AGAGAAGAGGAAAGAAAGAAAGCGGAAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 290
Qy 384 CGCACTAAGTGTCTATGATGAAGATTGAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 443
Db 291 GAGGAACCAAGAAAG 350
Qy 444 AGAAAAATGAACGGATAAAGCCACTTAAGCAATTCGAGAGAGAGAGAGAGAGAGAGAGAGAG 503
Db 351 AGAAAG 409
Qy 504 AGCTGAGGAGCAGAAAAAATTCAG 563
Db 410 AARAGAGAGGAG 463
Qy 564 TAAAG 623
Db 464 AAG 523
Qy 624 TCTTGCTCCAAAAGGGAGAAATTCAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 680
Db 524 AGATGAG 580

RESULT 10
US-09-270-767-20343
; Sequence 20343, Application US/09270767
; Patent No. 6703491
; GENERAL INFORMATION:
; APPLICANT: Homburger et al.
; TITLE OF INVENTION: Nucleic acids and proteins of *Drosophila melanogaster*
; FILE REFERENCE: File Reference: 7326-094
; CURRENT APPLICATION NUMBER: US/09/270,767
; CURRENT FILING DATE: 1999-03-17
; NUMBER OF SEQ ID NOS: 62517
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 20343
; LENGTH: 705
; TYPE: DNA
; ORGANISM: *Drosophila melanogaster*
US-09-270-767-20343

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; MOLECULE TYPE: oligodeoxynucleotide
; HYPOTHETICAL: NO
; ANTI-SENSE: YES
US-08-628-417-6

Query Match          5.7%; Score 72.4; DB 1; Length 240;
Best Local Similarity 57.5%; Pred. No. 2.2e-08;
Matches 130; Conservative 0; Mismatches 96; Indels 0; Gaps 0;

Qy 370 GAAGATCAAAATACGCACTAAGTGTGTCATGATGAAGATTGAAGAAGAGAGAGAAAA 429
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 11 GACAATAAACTTTAGAAATAATTTTACTAAAAAATAAAAAAAAAAAAAAAAAAAAA 70

Qy 430 ATAGAAGTATGAAGAAGAAATGACCGTAAAGCCCTAAAGCATTTGGAAGCTGGAAA 489
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 71 AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 130

Qy 490 GAATATCAAGAAAGCTGAGGAGCAAAAAAAAAAATTCAGAGAGAGAGAAATTTATGTCAA 549
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 131 AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 190

Qy 550 AAGANAAGCAATTAAGAAGAGAGAGAAAAAATAAATAATATAAGA 595
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 191 AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 236

RESULT 12
US-09-640-173-53/c
; Sequence 53, Application US/09640173
; Patent No. 6613515
; GENERAL INFORMATION:
; APPLICANT: Xu, Jiangchun
; APPLICANT: Stoik, John A.
; TITLE OF INVENTION: OVARIAN TUMOR SEQUENCES AND
; TITLE OF INVENTION: METHODS OF USE THEREFOR
; FILE REFERENCE: 210121.484C2
; CURRENT APPLICATION NUMBER: US/09/640.173
; CURRENT FILING DATE: 2000-08-15
; NUMBER OF SEQ ID NOS: 196
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 53
; LENGTH: 396
; TYPE: DNA
; ORGANISM: Homo sapien
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(396)
; OTHER INFORMATION: n = A,T,C or G
US-09-640-173-53

Query Match          5.7%; Score 72; DB 4; Length 396;
Best Local Similarity 54.9%; Pred. No. 3.4e-08;
Matches 135; Conservative 0; Mismatches 111; Indels 0; Gaps 0;

Qy 363 AAAGCGGAAGATCAAAAATACGCACCTAAGTGTGTCATGATGAAGATTGAAGAAGAGAG 422
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 247 ANAAGGNAANNAANNAANNTAANNAANNAANNAANNAANNAANNAANNAANNA 188

Qy 423 GAAAAAATGAAGATATGAAGAAGAAATGAACGGATTAAGCCCTAAAGCATTTGGAAGC 482
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 187 AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 128

Qy 483 CTGGAAGAATATCAAGAAGAAAGCTGAGGAGCAAAAAAAAAAATTCAGAGAGAGAGAAAT 542
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 127 AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 68

Qy 543 ATGTCAAAAAGAAAGCAAAATTTAAAGAAAGGAAGAAAAAATAAATATAGAGCTTTAC 602
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Db 67 AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 8

Qy 603 TAGAAA 608
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Db 7 AAAAAA 2

RESULT 13
US-09-713-550-53/c
; Sequence 53, Application US/09713550
; Patent No. 6617109
; GENERAL INFORMATION:
; APPLICANT: Xu, Jiangchun
; APPLICANT: Stoik, John A.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE
; TITLE OF INVENTION: THERAPY AND DIAGNOSIS OF OVARIAN CANCER
; FILE REFERENCE: 210121.484C4
; CURRENT APPLICATION NUMBER: US/09/713.550
; CURRENT FILING DATE: 2000-11-14
; NUMBER OF SEQ ID NOS: 205
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 53
; LENGTH: 396
; TYPE: DNA
; ORGANISM: Homo sapien
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(396)
; OTHER INFORMATION: n = A,T,C or G
US-09-713-550-53

Query Match          5.7%; Score 72; DB 4; Length 396;
Best Local Similarity 54.9%; Pred. No. 3.4e-08;
Matches 135; Conservative 0; Mismatches 111; Indels 0; Gaps 0;

Qy 363 AAAGCGGAAGATCAAAAATACGCACCTAAGTGTGTCATGATGAAGATTGAAGAAGAGAG 422
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Db 247 ANAAGGNAANNAANNAANNTAANNAANNAANNAANNAANNAANNAANNAANNA 188

Qy 423 GAAAAAATGAAGATATGAAGAAGAAATGAACGGATTAAGCCCTAAAGCATTTGGAAGC 482
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 187 AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 128

Qy 483 CTGGAAGAATATCAAGAAGAAAGCTGAGGAGCAAAAAAAAAAATTCAGAGAGAGAGAAAT 542
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 127 AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 68

Qy 543 ATGTCAAAAAGAAAGCAAAATTTAAAGAAAGGAAGAAAAAATAAATATAGAGCTTTAC 602
    ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 67 AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA 8

Qy 603 TAGAAA 608
    |||||
Db 7 AAAAAA 2

RESULT 14
US-09-825-294-53/c
; Sequence 53, Application US/09825294
; Patent No. 6710170
; GENERAL INFORMATION:
; APPLICANT: Xu, Jiangchun
; APPLICANT: Stoik, John A.
; APPLICANT: Algate, Paul A.
; APPLICANT: Fling, Steven P.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE
; TITLE OF INVENTION: THERAPY AND DIAGNOSIS OF OVARIAN CANCER
; FILE REFERENCE: 210121.484C5
; CURRENT APPLICATION NUMBER: US/09/825.294
; CURRENT FILING DATE: 2001-04-03
; NUMBER OF SEQ ID NOS: 215
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 53
; LENGTH: 396
; TYPE: DNA
; ORGANISM: Homo sapien
; FEATURE:
; NAME/KEY: misc_feature
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; LOCATION: (1)...(396)
; OTHER INFORMATION: n = A,T,C or G
; US-09-825-294-53

Query Match          5.7%; Score 72; DB 4; Length 396;
Best Local Similarity 54.9%; Pred. No. 3.4e-08;
Matches 135; Conservative 0; Mismatches 111; Indels 0; Gaps 0;

Qy 363 AAAGCGGAAGATCAAAAATACGCACCTAAAGTGTCAAGATTGAGAGAGAAGAGAG 422
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 247 ANAAGGNAANANANANANANNTTAAANANANANANANANANANANANANANAN 188
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Qy 423 GAAAAAATAGAAGATATGAAGAGAAATGAACGGATAAAGCCACTTAAGCAATTGGAAGC 482
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 187 AAAAAAANANANANANANANANANANANANANANANANANANANANANANANAN 128
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Qy 483 CTGGAAGATATCAAGAGAAAGCTCAGGAGCAAAAAAATTCAGAGAGAGAGAAATT 542
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 127 AAAAAAANANANANANANANANANANANANANANANANANANANANANANANAN 68
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Qy 543 ATGTCAAAAAGAAAGCAAAATTTAAAGAGAGGAAGAAAAAATAAATATTAAGAGTCTTAC 602
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 67 AAAAAAANANANANANANANANANANANANANANANANANANANANANANANAN 8
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Qy 603 TAGAAA 608
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 7 AAAAAA 2
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Search completed: July 8, 2005, 04:50:19
Job time : 248 secs

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RESULT 15
US-09-970-966-53/c
; Sequence 53, Application US/09970966
; Patent No. 6720146
; GENERAL INFORMATION:
; APPLICANT: Stolk, John A.
; APPLICANT: Molesh, David Alan
; APPLICANT: Fling, Steven P.
; APPLICANT: Xu, Jiangchun
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
; TITLE OF INVENTION: AND DIAGNOSIS OF OVARIAN CANCER
; FILE REFERENCE: 210121.484C6
; CURRENT APPLICATION NUMBER: US/09/970,966
; CURRENT FILING DATE: 2001-10-02
; NUMBER OF SEQ ID NOS: 215
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 53
; LENGTH: 396
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: 224, 225, 228, 235, 240, 246, 257, 266, 274, 279, 281, 282,
; LOCATION: 283, 285, 287, 288, 290, 291, 292, 293, 294, 295, 296, 297,
; LOCATION: 300, 301, 303, 307, 311, 313, 314, 317, 318, 319, 320, 321,
; LOCATION: 323, 324, 328, 329, 330, 336, 337, 338, 339, 340, 341
; OTHER INFORMATION: n = A,T,C or G
; NAME/KEY: misc feature
; LOCATION: 342, 343, 344, 345, 346, 347, 348, 349, 350, 351, 352, 356,
; LOCATION: 357, 358, 359, 362, 363, 364, 365, 366, 367, 373, 380, 381,
; LOCATION: 382, 385, 387, 388, 389, 390, 392
; OTHER INFORMATION: n = A,T,C or G
;
US-09-970-966-53

Query Match 5.7%; Score 72; DB 4; Length 396;
Best Local Similarity 54.9%; Pred. No. 3.4e-08;
Matches 135; Conservative 0; Mismatches 111; Indels 0; Gaps 0;

QY 363 AAAGCGGGAGATCAAAAATACGCACCTAAGTGTCTATGATGAAGATTGAAGAGAGAGAG 422
Db 247 ANAAGCNAAAANAAANAAANNTAAAAAANAAAAAANAAAAAANAAAAAANAAAAA 188
QY 423 GAAAAAATAGAGATATGAAGAAATGAACGGATAAAGCCACTAAAGCATTGGAAGC 482

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QY 961 AATAAGATGCCACTATTGTATTGTAACCGGGCTGCTGGCCACCTTAAACCTTAAACCTTA 1020
Db 1329 AATAAGATGCCACTATTGTATTGTAACCGGGCTGCTGGCCACCTTAAACCTTAAACCTTA 1388
QY 1021 CACAAGCTATTGAAGATTCTTCTAAGGCACTGGAATATTGATGCCACCTGTTTACAGAC 1080
Db 1389 CACAAGCTATTGAGATTCTTCTAAGGCACTGGAATATTGATGCCACCTGTTTACAGAC 1448
QY 1081 AATGCTAATCAAGAATGAAGGCACATGTACGACGTGGAAACAGCATTTCTGTCAACTAGAA 1140
Db 1449 AATGCTAATCAAGAATGAAGGCACATGTACGACGTGGAAACAGCATTTCTGTCAACTAGAA 1508
QY 1141 TTGTATGTAGAGCCCTACAGGATTATGAAGCGGCACTTAAGATTGATCCATCAACAAA 1200
Db 1509 TTGTATGTAGAGCCCTACAGGATTATGAAGCGGCACTTAAGATTGATCCATCAACAAA 1568
QY 1201 ATTGTACAAATTGATGCTGAGAAGATTTCGGAATGTAATTCAGGAAACAGAACTAAATCT 1260
Db 1569 ATTGTACAAATTGATGCTGAGAAGATTTCGGAATGTAATTCAGGAAACAGAACTAAATCT 1628
QY 1261 TAA 1263
Db 1629 TAA 1631
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RESULT 4

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US-10-681-199-2
; Sequence 2, Application US/10681199
; Publication No. US20040138441A1
; GENERAL INFORMATION:
; APPLICANT: KERE, Juha
; TITLE OF INVENTION: NOVEL HUMAN GENE FUNCTIONALLY RELATED TO DYSLEXIA
; FILE REFERENCE: 0933-0214P
; CURRENT APPLICATION NUMBER: US/10/681.199
; NUMBER OF SEQ ID NOS: 42
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 2
; LENGTH: 1993
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (369)..(1628)
; FEATURE:
; OTHER INFORMATION: human DYX1 mRNA as cDNA
US-10-681-199-2
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Query Match 100.0%; Score 1263; DB 19; Length 1993;
Best Local Similarity 100.0%; Pred. No. 5.9e-253;
Matches 1263; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGCTCTTTCAGGTTAGCGATTACAGCTGGCAGCAGACGAGACTGCGGTCTTTCTGTCT 60
Db 369 ATGCTCTTTCAGGTTAGCGATTACAGCTGGCAGCAGACGAGACTGCGGTCTTTCTGTCT 428
QY 61 CTGCCCCCTCAAGCGGTGTGCGTCAGACACGCGGTGTTCTGCAACGAAACCTATCTG 120
Db 429 CTGCCCCCTCAAGCGGTGTGCGTCAGACACGCGGTGTTCTGCAACGAAACCTATCTG 488
QY 121 AAGTCAACTTCTCCATTTTATTTTATTTAGGCAATTTCTTATGCTCCCATAGACGATGAG 180
Db 489 AAGTCAACTTCTCCATTTTATTTTATTTAGGCAATTTCTTATGCTCCCATAGACGATGAG 548
QY 181 AGCAGCAAGCAAGATTGGGAATGACACCATTTCTTACCTTGTATATAAAAGAGAGCG 240
Db 549 AGCAGCAAGCAAGATTGGGAATGACACCATTTCTTACCTTGTATATAAAAGAGAGCG 608
QY 241 GCCATGTGGGAGACCCCTTTTCTGTGACCGGGTGTTCACAAAGAGATGATGCAAGAAATAGA 300
Db 609 GCCATGTGGGAGACCCCTTTTCTGTGACCGGGTGTTCACAAAGAGATGATGCAAGAAATAGA 668
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RESULT 5

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US-10-681-199-19
; Sequence 19, Application US/10681199
; Publication No. US20040138441A1
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QY 301 GAAAAATCTATTTTCAAGCAAGAGAGAGCAAGAAAGAGCTTACAGAAAGCAAAAGCTGCA 360
Db 669 GAAAAATCTATTTTCAAGCAAGAGAGAGCAAGAAAGAGCTTACAGAAAGCAAAAGCTGCA 728
QY 361 GCAAAGCGGGAAGATCAAAAATACGCACTAAGTGTCTATGATGAAGATTGAAGAAGAGAG 420
Db 729 GCAAAGCGGGAAGATCAAAAATACGCACTAAGTGTCTATGATGAAGATTGAAGAAGAGAG 788
QY 421 AGCAAAAAATACAGATATGAAGAAATGAACGGATATAAAGCCATTAAGCATTTGGAA 480
Db 789 AGCAAAAAATACAGATATGAAGAAATGAACGGATATAAAGCCATTAAGCATTTGGAA 848
QY 481 GCCTGGAAGAAATATCAAAAGAAAGCTGAGAGCAAAAAAATTTTCAAGAGAGAGAGAA 540
Db 849 GCCTGGAAGAAATATCAAAAGAAAGCTGAGAGCAAAAAAATTTTCAAGAGAGAGAGAA 908
QY 541 TTATGTCAAAAGAAAGCAAAATTAAGAAAGAAAGAAAGAAATTAATAATTAAGAGTCTT 600
Db 909 TTATGTCAAAAGAAAGCAAAATTAAGAAAGAAAGAAAGAAATTAATAATTAAGAGTCTT 968
QY 601 ACTAGAAATTTGGCATCTAGAAATCTTGTCCAAAGGGAGAAATTCAGAAATATATTT 660
Db 969 ACTAGAAATTTGGCATCTAGAAATCTTGTCCAAAGGGAGAAATTCAGAAATATATTT 1028
QY 661 ACTGAGAAAGTTAAAGGAAGACAGTATTCTGCTCCTCGCTCTCTGTCAGTATTAAATC 720
Db 1029 ACTGAGAAAGTTAAAGGAAGACAGTATTCTGCTCCTCGCTCTCTGTCAGTATTAAATC 1088
QY 721 AACTTTACCCCTCGAGTATTCCCAACAGCTCTTTCGTTGTAATCAAGATGAGAGAGAG 780
Db 1089 AACTTTACCCCTCGAGTATTCCCAACAGCTCTTTCGTTGTAATCAAGATGAGAGAGAG 1148
QY 781 GAGTGCCTACACAAACAGCTGAGCGACGAGAGCAATGAATCTGCATAGCTGACTT 840
Db 1149 GAGTGCCTACACAAACAGCTGAGCGACGAGAGCAATGAATCTGCATAGCTGACTT 1208
QY 841 TGGCATTTAAAGAAAGAAAGAAAGAAACCCAGAAATGGTTGAAGGATAAAGGAAACAAATG 900
Db 1209 TGGCATTTAAAGAAAGAAAGAAAGAAACCCAGAAATGGTTGAAGGATAAAGGAAACAAATG 1268
QY 901 TTTGCAACGGAAACCTATTGTCAGCTATCAATGCATATATAATTTAGCCATAAGACTAAAT 960
Db 1269 TTTGCAACGGAAACCTATTGTCAGCTATCAATGCATATATAATTTAGCCATAAGACTAAAT 1328
QY 961 AATAAGATGCCACTATTGTATTGTAACCGGGCTGCTGGCCACCTTAAACCTTAAACCTTA 1020
Db 1329 AATAAGATGCCACTATTGTATTGTAACCGGGCTGCTGGCCACCTTAAACCTTAAACCTTA 1388
QY 1021 CACAAGGCTATTGAAGATTTCTTCTAAGGCACTGGAATTTATGATGCCACCTGTTTACAGAC 1080
Db 1389 CACAAGGCTATTGAAGATTTCTTCTAAGGCACTGGAATTTATGATGCCACCTGTTTACAGAC 1448
QY 1081 AATGCTAATCAAGAATGAAGGCACATGTACGACGTGGAAACAGCATTTCTGTCAACTAGAA 1140
Db 1449 AATGCTAATCAAGAATGAAGGCACATGTACGACGTGGAAACAGCATTTCTGTCAACTAGAA 1508
QY 1141 TTGTATGTAGAGCCCTACAGGATTATGAAGCGGCACTTAAGATTGATCCATCAACAAA 1200
Db 1509 TTGTATGTAGAGCCCTACAGGATTATGAAGCGGCACTTAAGATTGATCCATCAACAAA 1568
QY 1201 ATTGTACAAATTTGATGCTGAGAAGATTTCGGAATGTAATTCAGGAAACAGAACTAAATCT 1260
Db 1569 ATTGTACAAATTTGATGCTGAGAAGATTTCGGAATGTAATTCAGGAAACAGAACTAAATCT 1628
QY 1261 TAA 1263
Db 1629 TAA 1631
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; GENERAL INFORMATION:
; APPLICANT: KERE, Juha
; TITLE OF INVENTION: NOVEL HUMAN GENE FUNCTIONALLY RELATED TO DYSLEXIA
; FILE REFERENCE: 0933-0214P
; CURRENT APPLICATION NUMBER: US/10/681,199
; CURRENT FILING DATE: 2003-10-09
; NUMBER OF SEQ ID NOS: 42
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 19
; LENGTH: 1263
; TYPE: DNA
; ORGANISM: Pan paniscus
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (1)..(1260)
; US-10-681-199-19

Query Match          99.2%; Score 1253.4; DB 19; Length 1263;
Best Local Similarity 99.5%; Pred. No. 5e-251;
Matches 1257; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

Qy 1 ATGCTCTTTCAGGTTAGCGATTACAGCTGGCAGCAGACGAGACTGCGGTCTTCTGTCT 60
Db 1 ATGCTCTTTCAGGTTAGCGATTACAGCTGGCAGCAGACGAGACTGCGGTCTTCTGTCT 60

Qy 61 CTGCCCCCTCAAGGCGTGTGCTCAGACACGAGCGTGTCTGACCGGAAACTATCTG 120
Db 61 CTGCCCCCTCAAGGCGTGTGCTCAGACACGAGCGTGTCTGACCGGAAACTATCTG 120

Qy 121 AAGGTCACCTTTCCTCCATTTTATTTGAGGCACTTCTTATGCTCCCATGACGATGAG 180
Db 121 AAGGTCACCTTTCCTCCATTTTATTTGAGGCACTTCTTATGCTCCCATGACGATGAG 180

Qy 181 AGCAGCAAGCAAGATTGGGAATGACACCATTTGCTTTCACCTTGTATAAAAAAGAGCG 240
Db 181 AGCAGCAAGCAAGATTGGGAATGACACCATTTGCTTTCACCTTGTATAAAAAAGAGCG 240

Qy 241 GCCATGTGGGAGACCCCTTCTGTGTGACGGGTGTGACAAAGAGATGATCAAGAAATTAGA 300
Db 241 GCCATGTGGGAGACCCCTTCTGTGTGACGGGTGTGACAAAGAGATGATCAAGAAATTAGA 300

Qy 301 GAAATAATCTATTTTACAGCACAAGAGAGAGCAAAAGAGCTACAGAGCAAAAGCTGCA 360
Db 301 GAAATAATCTATTTTACAGCACAAGAGAGAGCAAAAGAGCTACAGAGCAAAAGCTGCA 360

Qy 361 GCAAGCGGGAAGATCAAAAATACCCACTAAGTGTGATGATGAAGATTGAAGAGAGAG 420
Db 361 GCAAGCGGGAAGATCAAAAATATGCACTAAGTGTGATGATGAAGATTGAAGAGAGAG 420

Qy 421 AGGAAAAAATAGAGAGATATGAAAGAAATGAAACGATATAAGCCACTAAAGCAATTGAA 480
Db 421 AGGAAAAAATAGAGAGATATGAAAGAAATGAAACGATATAAGCCACTAAAGCAATTGAA 480

Qy 481 GCCTGGAAGATATCAAGAGAAAGCTGAGGAGCAAAAGAAATTCAGAGAGAGAGAAA 540
Db 481 GCCTGGAAGATATCAAGAGAAAGCTGAGGAGCAAAAGAAATTCAGAGAGAGAGAAA 540

Qy 541 TTATGTCAAAAGAAAGCAAAATTAAGAGCAAAAGAAATTAATAATAAGAGTCTT 600
Db 541 TTATGTCAAAAGAAAGCAAAATTAAGAGCAAAAGAAATTAATAATAAGAGTCTT 600

Qy 601 ACTAGAAAATTTGGCATCTAGAAATCTTGCTCCAAAGGGAGAAATTCAGAAAAATATATT 660
Db 601 ACTAGAAAATTTGGCATCTAGAAATCTTGCTCCAAAGGGAGAAATTCAGAAAAATATATT 660

Qy 661 ACTGAGAGATTAAAGGAGACAGTATTCCTGCTCCTCGCTCTGTTGGCAGTATTAATATC 720
Db 661 ACTGAGAGATTAAAGGAGACAGTATTCCTGCTCCTCGCTCTGTTGGCAGTATTAATATC 720

Qy 721 AACTTTACCCCTCGAGTATTCACACAGCTCTTCTGTAATCACAGTAGCAGAGAGAG 780
Db 721 AACTTTACCCCTCGAGTATTCACACAGCTCTTCTGTAATCACAGTAGCAGAGAGAG 780
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Qy 781 GAGTGGCTACACAAACAGCTGAGCAGCAAGCAATGAATACTGACATAGCTGAACCTT 840
Db 781 GAGTGGCTGCAACAAACAGCTGAGCAGCAAGCAATGAATACTGACATAGCTGAACCTT 840

Qy 841 TGCATTTTAAAGAGAGAAAGAAACCCAGATGTTTGAAGGATAAAGGAAACAAATTTG 900
Db 841 TGCATTTTAAAGAGAGAGAAAGAAACCCAGATGTTTGAAGGATAAAGGAAACAAATTTG 900

Qy 901 TTTGCAACGGAAAACTATTTGGCAGCTATCAATGTCATATAATTTAGCCATAGACTAAAT 960
Db 901 TTTGCAACGAGAAAACTATTTGGCAGCTATCAATGTCATATAATTTAGCCATAGACTAAAT 960

Qy 961 AATAAGATGCCACTATTTGATTTGACCGGCTGCTGCGCCACTTAAACTTAAACCTTA 1020
Db 961 AATAAGATGCCACTATTTGATTTGACCGGCTGCTGCGCCACTTAAACTTAAACCTTA 1020

Qy 1021 CACAAGGCTATTGAAGATTCTTTAAGGCATCTGGAATTTATGATGCCACCTGTTACAGAC 1080
Db 1021 CACAAGGCTATTGAAGATTCTTTAAGGCATCTGGAATTTATGATGCCACCTGTTACAGAC 1080

Qy 1081 AATGCTAATCAAGAAATGAAGCACAATGACGCTGGAACAGCATTCTGTCAACTAGAA 1140
Db 1081 AATGCTAATCAAGAAATGAAGCACAATGACGCTGGAACAGCATTCTGTCAACTAGAA 1140

Qy 1141 TTGTATGTAGAGGCTTACAGGATTTGAGGCGCACTTAAGATTGATCCCAACAA 1200
Db 1141 TTGTATGTAGAGGCTTACAGGATTTGAGGCGCACTTAAGATTGATCCCAACAA 1200

Qy 1201 ATTGTACAAATTTGATGCTGAGAAGATTCCGGAATGTAATTTCAAGGAAACAGAACT 1260
Db 1201 ATTGTACAAATTTGATGCTGAGAAGATTCCGGAATGTAATTTCAAGGAAACAGAACT 1260

Qy 1261 TAA 1263
Db 1261 TAA 1263

RESULT 6
US-10-681-199-13
; Sequence 13, Application US/10681199
; Publication No. US20040138441A1
; GENERAL INFORMATION:
; APPLICANT: KERE, Juha
; TITLE OF INVENTION: NOVEL HUMAN GENE FUNCTIONALLY RELATED TO DYSLEXIA
; FILE REFERENCE: 0933-0214P
; CURRENT APPLICATION NUMBER: US/10/681,199
; CURRENT FILING DATE: 2003-10-09
; NUMBER OF SEQ ID NOS: 42
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 13
; LENGTH: 1263
; TYPE: DNA
; ORGANISM: Pan troglodytes
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (1)..(1260)
; US-10-681-199-13

Query Match          99.1%; Score 1251.8; DB 19; Length 1263;
Best Local Similarity 99.4%; Pred. No. 1.1e-250;
Matches 1256; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

Qy 1 ATGCTCTTTCAGGTTAGCGATTACAGCTGGCAGCAGACGAGACTGCGGTCTTCTGTCT 60
Db 1 ATGCTCTTTCAGGTTAGCGATTACAGCTGGCAGCAGACGAGACTGCGGTCTTCTGTCT 60

Qy 61 CTGCCCCCTCAAGGCGTGTGCTCAGACACGAGCGTGTTCGACGGAAACTATCTG 120
Db 61 CTGCCCCCTCAAGGCGTGTGCTCAGACACGAGCGTGTTCGACGGAAACTATCTG 120

Qy 121 AAGTCAACTTTCCTCCATTTTATTTGAGGCACTTCTTATGCTCCCATGACGATGAG 180
Db 121 AAGTCAACTTTCCTCCATTTTATTTGAGGCACTTCTTATGCTCCCATGACGATGAG 180
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QY 661 ACTGAGAGTTAAAGGAAGACAGTATTCCTGCTCCTCGCTCTGTTGGCAGTATTTAAATC 720
DB |||||
DB 661 ACTGAGAGTTAAAGGAAGACAGTATTCCTGCTCCTCGCTCTGTTGGCAGTATTTAAATC 720
QY 721 AACTTTACCCCTCGAGTATTCCTCAACAGCTCTTCGTGATCAACAGTACGAGAGGAG 780
DB |||||
DB 721 AACTTTACCCCTCGAGTATTCCTCAACAGCTCTTCGTGATCAACAGTACGAGAGGAG 780
QY 781 GAGTGGCTACACAAACAGCTGAGGACGAGAGCAATGAATPACTGACATAGCTGAAT 840
DB |||||
DB 781 GAGTGGCTACACAAACAGCTGAGGACGAGAGCAATGAATPACTGACATAGCTGAAT 840
QY 841 TCGGATTTAAAGGAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAG 900
DB |||||
DB 841 TCGGATTTAAAGGAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAG 900
QY 901 TTTGCAACGAGAAACTATTTGACAGCTATCAATGCATATAATTTAGCCATAGACTAAAT 960
DB |||||
DB 901 TTTGCAACGAGAAACTATTTGACAGCTATCAATGCATATAATTTAGCCATAGACTAAAT 960
QY 961 AATAAGTGCCTACTATTTGAAACCGGGCTGCTGCACTTAAACTTAAACTTAA 1020
DB |||||
DB 961 AATAAGTGCCTACTATTTGAAACCGGGCTGCTGCACTTAAACTTAAACTTAA 1020
QY 1021 CACAAGGCTATTGAAGATTTCTTAAGGCACTGGAATTTATGATGCCACCTGTTACAGAC 1080
DB |||||
DB 1021 CACAAGGCTATTGAAGATTTCTTAAGGCACTGGAATTTATGATGCCACCTGTTACAGAC 1080
QY 1081 AATGCTAATCAAGAAATGAAGCACTATGACGCTGGAACAGCACTTCTGCACTAGAA 1140
DB |||||
DB 1081 AATGCTAATCAAGAAATGAAGCACTATGACGCTGGAACAGCACTTCTGCACTAGAA 1140
QY 1141 TTGATGATGAGAGGCTACAGGATTTGAAGCGGCACTTAAGATTTGATCCATCAACAA 1200
DB |||||
DB 1141 TTGATGATGAGAGGCTACAGGATTTGAAGCGGCACTTAAGATTTGATCCATCAACAA 1200
QY 1201 ATTGACAAATGATGCTGAGAGATTCGGAAATGTAATCAAGGAACAGAACTTAAATCT 1260
DB |||||
DB 1201 ATTGACAAATGATGCTGAGAGATTCGGAAATGTAATCAAGGAACAGAACTTAAATCT 1260
QY 1261 TAA 1263
DB |||||
DB 1261 TAA 1263

RESULT 8

US-10-681-199-17
; Sequence 17, Application US/10681199
; Publication No. US20040138441A1
; GENERAL INFORMATION:
; APPLICANT: KERE, Juha
; TITLE OF INVENTION: NOVEL HUMAN GENE FUNCTIONALLY RELATED TO DYSLEXIA
; FILE REFERENCE: 0933-0214P
; CURRENT APPLICATION NUMBER: US/10/681,199
; CURRENT FILING DATE: 2003-10-09
; NUMBER OF SEQ ID NOS: 42
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 17
; LENGTH: 1263
; TYPE: DNA
; ORGANISM: Pongo pygmaeus
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (1)..(1260)
US-10-681-199-17

Query Match 98.5%; Score 1243.8; DB 19; Length 1263;
Best Local Similarity 99.0%; Pred. No. 5e-249;
Matches 1251; Conservative 0; Mismatches 12; Indels 0; Gaps 0;

QY 1 ATGCTCTTCAGGTTAGGATTTACAGCTGGCAGCAGACGAGACTGCGGCTCTTCTGTCT 60
|||||

DB 1 ATGCTCTTCAGGTTAGGATTTACAGCTGGCAGCAGACGAGACTGCGGCTCTTCTGTCT 60
QY 61 CTGCCCCCTCAAAGGCGTGTGCGTCAGAGACACGACGCTGTTCTGCAACGAAAACTATCTG 120
DB |||||
DB 61 CTGCCCCCTCAAAGGCGTGTGCGTCAGAGACACGACGCTGTTCTGCAACGAAAACTATCTG 120
QY 121 AAGTCACTTTCCTCCATTTTATTTGAGGCAATTTCTTTATGCTCCCATAGACGATGAG 180
DB |||||
DB 121 AAGTCACTTTCCTCCATTTTATTTGAGGCAATTTCTTTATGCTCCCATAGACGATGAG 180
QY 181 AGCAGCAAGCAAGATTTGGGAATGACACCATTTGCTTTCACCTTGTATATAAAAAAGAGCG 240
DB |||||
DB 181 AGCAGCAAGCAAGATTTGGGAATGACACCATTTGCTTTCACCTTGTATATAAAAAAGAGCG 240
QY 241 GCCATGTCGGAGACCTTTCTGTGACGGGTGTGAACAAAGAGATGATGCAAAAGATTTAGA 300
DB |||||
DB 241 GCCATGTCGGAGACCTTTCTGTGACGGGTGTGTGACAAAGAGACGATGCAAAAGATTTAGA 300
QY 301 GAAAAATCTATTTTACAAGCAACAGAGAGCAAAAGAGCTTACAGAGCAAAAGCTGCA 360
DB |||||
DB 301 GAAAAATCTATTTTACAAGCAACAGAGAGCAAAAGAGCTTACAGAGCAAAAGCTGCA 360
QY 361 GCAAGCGGGAAGATCAAAAAATACGCACTAAGTGTGATGAAGATTTGAAGAGAGAG 420
DB |||||
DB 361 GCAAGCGGGAAGATCAAAAAATATGCACTAAGTGTGATGAAGATTTGAAGAGAGAG 420
QY 421 AGGAAAAAATAGAGATATGAAAAAATGAAACGGATTAAGAGCCACTTAAAGCAATTTGAA 480
DB |||||
DB 421 AGGAAAAAATAGAGATATGAAAAAATGAAACGGATTAAGAGCCACTTAAAGCAATTTGAA 480
QY 481 GCCTGGAAGAAATATCAAGAAAGAGCTGAGAGCAAAAAAATTTTCAAGAGAGAGAA 540
DB |||||
DB 481 GCCTGGAAGAAATATCAAGAAAGAGCTGAGAGCAAAAAAATTTTCAAGAGAGAGAA 540
QY 541 TTATGCTCAAAAGAAAGCAAAATTAAGAAAGAAAGAAAGAAAGAAATATAAGAGTCTT 600
DB |||||
DB 541 TTATGCTCAAAAGAAAGCAAAATTAAGAAAGAAAGAAAGAAAGAAATATAAGAGTCTT 600
QY 601 ACTGAAATTTGGCATCTAGAAATCTTCTCCAAAGGGAGAAATTTCAAGAAATATATTT 660
DB |||||
DB 601 ACTGAAATTTGGCATCTAGAAATCTTCTCCAAAGGGAGAAATTTCAAGAAATATATTT 660
QY 661 ACTGAGAAATTTAAGGAAGACAGTATTCCTGCTCCTCGCTCTGTTGGCAGTATTTAAATC 720
DB |||||
DB 661 ACTGAGAAATTTAAGGAAGACAGTATTCCTGCTCCTCGCTCTGTTGGCAGTATTTAAATC 720
QY 721 AACTTTACCCCTCGAGTATTCCTCAACAGCTCTTCGTGAATCAACAGTACGAGAGGAG 780
DB |||||
DB 721 AACTTTACCCCTCGAGTATTCCTCAACAGCTCTTCGTGAATCAACAGTACGAGAGGAG 780
QY 781 GAGTGGCTACACAAACAGCTGAGGACGAGAGCAATGAATPACTGACATAGCTGAAT 840
DB |||||
DB 781 GAGTGGCTACACAAACAGCTGAGGACGAGAGCAATGAATPACTGACATAGCTGAAT 840
QY 841 TCGGATTTAAAGGAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAG 900
DB |||||
DB 841 TCGGATTTAAAGGAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAGAAAG 900
QY 901 TTTGCAACGAGAAACTATTTGACAGCTATCAATGCATATAATTTAGCCATAGACTAAAT 960
DB |||||
DB 901 TTTGCAACGAGAAACTATTTGACAGCTATCAATGCATATAATTTAGCCATAGACTAAAT 960
QY 961 AATAAGTGCCTACTATTTGAAACCGGGCTGCTGCACTTAAACTTAAACTTAA 1020
DB |||||
DB 961 AATAAGTGCCTACTATTTGAAACCGGGCTGCTGCACTTAAACTTAAACTTAA 1020
QY 1021 CACAAGGCTATTGAAGATTTCTTAAGGCACTGGAATTTATGATGCCACCTGTTACAGAC 1080
DB |||||
DB 1021 CACAAGGCTATTGAAGATTTCTTAAGGCACTGGAATTTATGATGCCACCTGTTACAGAC 1080
QY 1081 AATGCTAATCAAGAAATGAAGCACTATGACGCTGGAACAGCACTTCTGCACTAGAA 1140
DB |||||
DB 1081 AATGCTAATCAAGAAATGAAGCACTATGACGCTGGAACAGCACTTCTGCACTAGAA 1140

QY 121 AAGTCAACTTCTCCCAATTTTATTGAGGCAATTTCTTTATGCTCCATAGACGATGAG 180
DB 168 AAGGTTAACTTCTCTCCCAATTTTATTGAGGCTGTTTCTATGCTCCATAGATGATGGG 227
QY 181 AGCAGCAAGCAAGATTTGGGAATGACACCATTTGCTTCCATGATATAAAGAGCGG 240
DB 228 AAGAGCAAGCCAGATTTGGAAATGACAGATTTCTTTTACATTTGATATAAAGAGGCCA 287
QY 241 GCCATGTGGGAGACCTTTTCTGTGACGGGTGTGTGACAAAGAGATGATCAAGAATTAGA 300
DB 288 GTTCTGTGGATAGCTTTTCTGTGCGGGTGTGTGATAAAGAGATGATGACAGATAAGA 347
QY 301 GAAAAATCTATTTCAGACACAGAGAGACGAAAAAGAGCTACAGAAAGCAAAAGCTGCA 360
DB 348 GAAAAATCTATTCTGCAAGCAGAGAAAGCAAAAGAGGCCACAGAAAGCAAAAGCTGTT 407
QY 361 GCAAGCGGGAAGATCAAAATACGCACTTAAGTGTATGATGAAGATTAAGAAAGAGAG 420
DB 408 GCCAGCGAGAGACACAGATACGCACTTAGCGGAGATGATGAAGATTAAGAAAGAGAG 467
QY 421 AGGAAAAAATAGAGATATGAAAGAAATGAACGGATAAAGCCACTAAAGCAATTTGAA 480
DB 468 AGGAAAAAATCGAGATCTGAAGAAATGAACGGAAAAAGGCAACTAGCGAATTAGAA 527
QY 481 GCCTGGAAGAAATATCAAGAAAGCTGAGGAGCAAAAAAATTCAGAGAGAGAGAAA 540
DB 528 GCGTGGAAAGAAATGTCAAAAGAAAGCTGACGCAAAAAAAGAGTCCAGAGGAAAGAGAA- 586
QY 541 TTATGTCAAAAGAAAAAGCAAAATTAAGAGAGAGAGAAAAAATAAATAAAGAGTCTT 600
DB 587 --ACCGCTCGAGGGAAGCAAGCT---GAAAGACCAAAAGCTCTAAACCTCGGGGTTG 641
QY 601 ACTAGAAATTTGGCATCTAGAAATCTTGCTCCAAAAGGGAGAAATTCAGAAAAATATTT 660
DB 642 CCCGGAAGGCCCACTCGCTCCGCAAGAGGGAGGAAATTTGGAAAAACATATTT 701
QY 661 ACTGAGAAATTAAGGAAGACGATATCTGCTCTCTGCTGCTGTTGGCAGTATTAAGATC 720
DB 702 CCTGAGAAATTAAGGAAGACAGAGTCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 761
QY 721 AACTTTACCTCCGAGTATCCCAACAGCTCTTGTGATCATCAGTAGCAGAGAGGAG 780
DB 762 AGCTTTACCTCCGAGTGTCCCAACAGCACTTTGGGAAATCCCAAGTCGCAAGAGAGAG 821
QY 781 GAGTGGCTACACAAAGAGCTGAGGACGAGCAAGCAATGAATATCTGATAGCTGAACCTT 840
DB 822 GAGTGGCTGCATTAACAGACAGAGACGAGAGGCCATGAGCACTGACCTTCTGAGTTC 881
QY 841 TGGATTTAAAGAGAAAGAAAGAACCCAGAAATGTTGAAAGGATAAAGGAAACAAATTTG 900
DB 882 TTTGACTTTAAAGAGAAAGAGAGGAATCCAGACTGGTTGAAAGACAAAGGAAACAAATTTG 941
QY 901 TTTGCAAGGAAACTATTTGGCAGCTATCAATGATATATTTAGCCATTAAGCACTAAT 960
DB 942 TTTGCAACAGAAAACTATTTGGCAGCGTTGATGATATATTTAGCCATTAAGCACTAAT 1001
QY 961 AATAAGATGCCATTTGTTGTTGAAACGGGCTGCTGCTGCACTTAAACTTAAAGAACTTTA 1020
DB 1002 TGTAAATGCCATTTATTTGTTGTTGAAATCGGGCTGCTGCTGCACTTAAACTTAAAGAACTTTA 1061
QY 1021 CACAAGGCTATTTGAGATTTCTTTAAGGCACCTGGAATTTATTTGATGCCACCTGTTTACAGAC 1080
DB 1062 CACAAGGCCATCGAGGACTCTTCTTAAGGCACCTAGAGTTTATTTGACACCACTGTTTGTCTGAC 1121
QY 1081 AATGCTAATGCAAGATGAGGCACATGTAAGAGTGAACAGCACTTCTGCTCACTAGAA 1140
DB 1122 AATGCTAATGCAAGATGAGGCACATGTAAGAGTGAACAGCACTTCTGCTCACTAGAA 1181
QY 1141 TTGATGTAGAGGCTACAGATTTATGAAGCGGCACTTAAGATTTGATCCATCCAAACAA 1200
DB 1182 TTGATGTAGAGGCTTCAAGATTTATGAAGCTGCACTTAAGATTTGATCCAGCCAGCCACACA 1241

QY 1201 ATTGTACAAATTTGATGCTGACAGATTCGGATGTAAATTCAGGACAGAACTAAATCT 1260
DB 1242 GTTGTACAGAACGATGCGAGAGATTCGGAATATATTAATTCAGGGACGCGACTGAAGTCT 1301

RESULT 11

US-10-681-199-4
; Sequence 4, Application US/10681199
; Publication No. US2004013841A1
; GENERAL INFORMATION:

; APPLICANT: KERE, Juha
; TITLE OF INVENTION: NOVEL HUMAN GENE FUNCTIONALLY RELATED TO DYSLEXIA
; FILE REFERENCE: 0933-0214P
; CURRENT APPLICATION NUMBER: US/10/681,199
; CURRENT FILING DATE: 2003-10-09
; NUMBER OF SEQ ID NOS: 42
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 4

; LENGTH: 1697

; TYPE: DNA

; ORGANISM: Mus musculus

; FEATURE:

; NAME/KEY: CDS

; LOCATION: (48)..(1307)

; FEATURE:

; OTHER INFORMATION: murine DYX1 mRNA as cDNA

US-10-681-199-4

Query Match 66.6%; Score 840.8; DB 19; Length 1697;

Best Local Similarity 80.3%; Pred. No. 4.7e-165; Indels 6; Gaps 2;

Matches 1012; Conservative 0; Mismatches 242;

QY 1 ATGCTCTTTCAGGTTAGCGATTACAGCTGGCAGCAGACGAAAGACTGCGGTCTTTCTGTCT 60

DB 48 ATGCAGGTGCGAGTGAGCGAATTCAGCTGGCAGCAGACGCGGCCACGATCTTCTGTGCG 107

QY 61 CTGCCCCCAAGGCGTGTGCTGAGACACGACGCTGTTCTGACCGGAAAACTATCTG 120

DB 108 CTGCTCTGCGGGGCGTCTGCGTGGCGATGCTGACGTAATCTGTGGGGAAGTTACCTG 167

QY 121 AAGTCAACTTCTCCATTTTATTTGAGGCAATTTCTTTATGCTCCCATAGACCATGAG 180

DB 168 AAGTTAACTTCTCTCCATTTTATTTGAGGCTGTTCTTATGCTCCCATAGATGAGG 227

QY 181 AGCAGCAAGCAAGATTTGGGAATGACACCAATTTGTTTACCTTTGTATATAAAGAAAGCG 240

DB 228 AAGAGCAAGCAAGATTTGGAAATGACACGATTTCTTTTCACTGTATAAAGAGGCCA 287

QY 241 GCCATGTGGGAGACCTTTCTGTGACGGGTGTTGACAAAGAGATGATCAAGAATTAGA 300

DB 288 GTTCTGTGGGATAGCTTTCTGTGCGGGTGTGATAAAGAGATGATGACAGAAATAAGA 347

QY 301 GAAAAATCTATTTTACAGCAAGAGAGAGCAAAAGAGCTACAGAAAGCAAAAGCTGCA 360

DB 348 GAAAAATCTATTTTACAGCAAGAGAGAGCAAAAGAGCTGCAAAAGAGCAAAAGCTGTT 407

QY 361 GCAAGCGGGAAGATCAAAATATAGCCATAGTGTCTATGATCAAGATTTGAAGAAGAGAG 420

DB 408 GCCAAGCGGGAAGACCAAGATAGCAGATAGCAGGATGATGAGATTTGAAGAAGAGAG 467

QY 421 AGGAAAAAATAGAAATATGAAAGAAATGAAAGGATAAAGCCACTTAAAGCATTTGAA 480

DB 468 AGGAAAAAATCTGAAGATCTGAAAGAAATGAAAGGAAAAAGGCAACTAGCGAATTAGAA 527

QY 481 GCCTGGAAGAAATCAAGAAAGAGCTGAGGAGCAAAAAAATTCAGAGAGAGAGAGAA 540

DB 528 GCGTGGAAAGAAATGTCAAAAGAAAGAGCTGACGCAAAAAAAGAGTCCAGAGAGAGAGAA- 586

QY 541 TTATGTCAAAAGAAAAAGCAAAATTAAGAGAGAGAGAAAAAATAAATAAAGAGTCTT 600

DB 587 --ACCGCTCGAGGGAAGCAAGCT---GAAAGACCAAAAGCTCTTAAACCTCGGGGTTG 641

QY 601 ACTAGAAATTTGGCATCTAGAAATCTTGTCTCCAAAGGGAGAAATTCAGAAAAATATATTT 660

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Db 642 CCCGGAAGGCCCCACCCTCGCTCCCAAGAGGGAGGAATTCGGAAACATATTT 701
Qy 661 ACTGAGAAGTAAAGGAAGACAGTATTCCTGCTCTCGCTCTGTGGCAGTATTAATC 720
Db 702 CCTGAGAAGTAAAGGAAGACAGAGTCCCTCGCTCGCTCGCTCGCTCGCTCGCTCG 761
Qy 721 AACTTTACCCCTCGAGTATTCACCAAGCTCTTGGTGAATCAACAAGTAGCAGAGAGGAG 780
Db 762 AGCTTTACCCCTCGAGTATTCACCAAGCTCTTGGGAAATCCCAAGTCGCAAGAGGAG 821
Qy 781 GAGTGGCTACACAAAAGCTGAGGCAAGAGCAATGAATCTGCATAGCTGAACCTT 840
Db 822 GAGTGGCTGCTAATAACCAAGCAGAGCACGGAGAGCCATGACCTTCCTGAGTTC 881
Qy 841 TGGGATTTAAAGAAAGAAAGAACCCAGATGGTGAAGAGTAAGAAAGAAACAATTG 900
Db 882 TTTGACTTAAAGAAAGAGAGAGGATCCAGACTGGTGAAGACACAAGGGAACAATTG 941
Qy 901 TTTGCAAGCAAGAACTATTTGGCAGCTATCAATGCATATATTTAGCCATAAGCAATAAT 960
Db 942 TTTGCAACAGAAACTATTTGGCAGCGGTGATGCATATATTTAGCCATACGACTGAAC 1001
Qy 961 AATAAGATGCCACTATTTGATTTGAACCGGCTGCTTGCACCTTAAACTAAAAACTTA 1020
Db 1002 TGTAAAGATCCATATTTGATTTGAATCGGCTGCTTGCACCTCAATTAATAAACTTA 1061
Qy 1021 CACAAGCTATTTGAAGATTTCTTAAGGCACTGGAATTTATGATGCCACCTGTTCAGAC 1080
Db 1062 CACAAGGCCATCGAGGACTCTTCTAAGGCCATAGAGTTATTTAGACACCCTGTTCGTGAC 1121
Qy 1081 AATGCTAATCAAGAAAGGACATGTAGGCTGGAACGATCTCTCAACTAGAA 1140
Db 1122 AATGCCAATCAAGAAAGGACACGCTCGAGGGACAGCGTCTGCTCAACTAGAA 1181
Qy 1141 TTGTATCTAGAAGCCCTACAGGATTTAAGAGCGGCACTTAAGATTTGATCCATCCAAACAA 1200
Db 1182 TTGTATCTAGAAGCCCTGCAAGATTTAAGAGCTGCACTTAAGATTTGACCCAGCCACACA 1241
Qy 1201 ATTGTACAAATTTGATCTGAGAGATTCGGAATGTAATTCAGGAACAGAACTAAATCT 1260
Db 1242 GTTGTACAGAACGATGCAGAGAGATTCGGAATATATTTCAAGGGACGGCACTGAAGTCT 1301
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RESULT 12

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US-10-242-535A-30001
; Sequence 30001, Application US/10242535A
; Publication No. US20040013663A1
; GENERAL INFORMATION:
; APPLICANT: ChondroGene Inc.
; APPLICANT: Liew, C.C.
; TITLE OF INVENTION: Compositions and Methods Relating to Osteoarthritis
; FILE REFERENCE: 4231/2005
; CURRENT APPLICATION NUMBER: US/10/242,535A
; CURRENT FILING DATE: 2002-09-12
; PRIOR APPLICATION NUMBER: US 10/085,783
; PRIOR FILING DATE: 2002-02-28
; PRIOR APPLICATION NUMBER: US 60/305,340
; PRIOR FILING DATE: 2001-07-13
; PRIOR APPLICATION NUMBER: US 60/275,017
; PRIOR FILING DATE: 2001-03-12
; PRIOR APPLICATION NUMBER: US 60/271,955
; PRIOR FILING DATE: 2001-02-28
; NUMBER OF SEQ ID NOS: 58994
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 30001
; LENGTH: 464
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (263)..(263)
; OTHER INFORMATION: n is a, c, g, or t
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; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (321)..(321)
; OTHER INFORMATION: n is a, c, g, or t
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (347)..(347)
; OTHER INFORMATION: n is a, c, g, or t
US-10-242-535A-30001
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Query Match 31.8%; Score 401.6; DB 17; Length 464;
Best Local Similarity 98.3%; Pred. No. 8.7e-74;
Matches 404; Conservative 0; Mismatches 7; Indels 0; Gaps 0;
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Qy 1 ATGCCTCTTCAGCTTAGCGATTACAGCTGGCAGCAGACGAAGACTGCGGTCTTTCTCTCT 60
Db 49 ATGCCTCTTCAGGTTAGCGATTACAGCTGGCAGCAGACGAAGACTGCGGTCTTTCTCTCT 108
Qy 61 CTGCCCCCTCAAGCGGTGTCGCTCAGAGACACGACCGTGTTCGACGGAATACTATCTG 120
Db 109 CTGCCCCCTCAAGCGGTGTCGCTCAGAGACACGACCGTGTTCGACGGAATACTATCTG 168
Qy 121 AAGGTCAACTTCTCTCAATTTTATTTAGGCAATTTCTTTATGCTCCCATAGAGATGAG 180
Db 169 AAGGTCAACTTCTCTCAATTTTATTTAGGCAATTTCTTTATGCTCCCATAGAGATGAG 228
Qy 181 AGCAGCAAGCAAGCAAGATTTGGGAATGACACCATTTGCTTACCTTGTATAAAAAAGAGCG 240
Db 229 AGCAGCAAGCAAGCAAGATTTGGGAATGACACCATTTGCTTACCTTGTATAAAAAAGAGCG 288
Qy 241 GCCATGTGGGAGACCCCTTTCTGTGACGGGTGTTGACAAAGAGATGATSCAAAGAAATTAGA 300
Db 289 GCCATGTGGGAGACCCCTTTCTGTGACGGGTGTTGACAAAGAGATGATSCAAAGAAATTANA 348
Qy 301 GAAAAATCTATTTTACAAGCACAAAGAGAGAGCAAAAAGAGCTACAGAAGCAAAAGCTGCA 360
Db 349 GAAAAATCTATTTTACAAGCACAAAGAGAGAGCAAAAAGAGCTACAGAAGCAAAAGCTGCA 408
Qy 361 GCAAGCGGGAAGATCAAAAATACGCACTAAGTGTCTATGATGAAGATTGAA 411
Db 409 GCAAGCGGGAAGATCAAAAATACGCACTAAGTGTCTATGATGAAGGGAGAA 459
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RESULT 13

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US-10-085-783A-30001
; Sequence 30001, Application US/10085783A
; Publication No. US20040037841A1
; GENERAL INFORMATION:
; APPLICANT: ChondroGene Inc.
; APPLICANT: Liew, C.C.
; TITLE OF INVENTION: Compositions and Methods Relating to Osteoarthritis
; FILE REFERENCE: 4231/2002
; CURRENT APPLICATION NUMBER: US/10/085,783A
; CURRENT FILING DATE: 2002-02-28
; PRIOR APPLICATION NUMBER: US 60/305,340
; PRIOR FILING DATE: 2001-07-13
; PRIOR APPLICATION NUMBER: US 60/275,017
; PRIOR FILING DATE: 2001-03-12
; PRIOR APPLICATION NUMBER: US 60/271,955
; PRIOR FILING DATE: 2001-02-28
; NUMBER OF SEQ ID NOS: 58994
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 30001
; LENGTH: 464
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (263)..(263)
; OTHER INFORMATION: n is a, c, g, or t
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (321)..(321)
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; OTHER INFORMATION: n is a, c, g, or t
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (347)..(347)
; OTHER INFORMATION: n is a, c, g, or t
US-10-085-783A-30001

Query Match      31.8%; Score 401.6; DB 18; Length 464;
Best Local Similarity 98.3%; Pred. No. 8.7e-74;
Matches 404; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 1 ATGCTCTTTACGTTAGCGATTACAGCTGGCAGCAGAGCAAGACTGCGTCTTTCTGCT 60
DB 49 ATGCTCTTTACGTTAGCGATTACAGCTGGCAGCAGAGCAAGACTGCGTCTTTCTGCT 108
QY 61 CTGCCCCCTCAAGGGCGTGTGCGTACAGACACGACGCTGTTCTGACCGAAACTATCTG 120
DB 109 CTGCCCCCTCAAGGGCGTGTGCGTACAGACACGACGCTGTTCTGACCGAAACTATCTG 168
QY 121 AGGTCAACTTTCCTCCATTTTATTTAGGCAATTTCTTATGCTCCCATAGACGATGAG 180
DB 169 AGGTCAACTTTCCTCCATTTTATTTAGGCAATTTCTTATGCTCCCATAGACGATGAG 228
QY 181 AGCAGCAAGCAAGATGGGAATGACACCATTTGCTTCACTCTGTATAAAAAAGACG 240
DB 229 AGCAGCAAGCAAGATGGGAATGACACCATTTGCTTCACTCTGTATAAAAAAGACG 288
QY 241 GCCATGTGGGAGACCTTTCTGTGACGGGTGTGTGACAAAGAGATGATGCAAGAAATTAGA 300
DB 289 GCCATGTGGGAGACCTTTCTGTGACGGGTGTGTGACAAAGAGATGATGCAAGAAATTANA 348
QY 301 GAAAAATCTATTTTACAGACCAAGAGAGAGCAAGAAAGAGCTACAGAGCAAAAGCTGCA 360
DB 349 GAAAAATCTATTTTACAGACCAAGAGAGAGCAAGAAAGAGCTACAGAGCAAAAGCTGCA 408
QY 361 GCAAGCGGAGAGATCAAAATACCACTAGTGTGATGATGCAAGATTTGAA 411
DB 409 GCAAGCGGAGAGATCAAAATACCACTAGTGTGATGATGCAAGAGGAGAA 459

RESULT 14
US-09-918-995-22675
; Sequence 22675, Application US/09918995
; Publication No. US20030073623A1
; GENERAL INFORMATION:
; APPLICANT: Hyseq, Inc.
; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
; FILE REFERENCE: 20411-756
; CURRENT APPLICATION NUMBER: US/09/918,995
; CURRENT FILING DATE: 2001-07-30
; PRIOR APPLICATION NUMBER: US/09/235,076
; NUMBER OF SEQ ID NOS: 38054
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 22675
; LENGTH: 488
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-918-995-22675

Query Match      29.3%; Score 369.8; DB 10; Length 488;
Best Local Similarity 99.5%; Pred. No. 3.7e-67;
Matches 371; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 35 AGACCAAGACTGCGGTCTTTCTGCTCTGCCCTCAAGGGCGTGTGCGTACAGACACGG 94
DB 28 AGACCAAGACTGCGGTCTTTCTGCTCTGCCCTCAAGGGCGTGTGCGTACAGACACGG 87
QY 95 ACGTGTCTGACGCGAAACTATCTGAAAGTCAACTTTCCTCCATTTTATTTAGGCAAT 154
DB 88 ACGTGTCTGACGCGAAACTATCTGAAAGTCAACTTTCCTCCATTTTATTTAGGCAAT 147

; OTHER INFORMATION: n is a, c, g, or t
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (347)..(347)
; OTHER INFORMATION: n is a, c, g, or t
US-10-085-783A-30001

Query Match      31.8%; Score 401.6; DB 18; Length 464;
Best Local Similarity 98.3%; Pred. No. 8.7e-74;
Matches 404; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

QY 1 ATGCTCTTTACGTTAGCGATTACAGCTGGCAGCAGAGCAAGACTGCGTCTTTCTGCT 60
DB 49 ATGCTCTTTACGTTAGCGATTACAGCTGGCAGCAGAGCAAGACTGCGTCTTTCTGCT 108
QY 61 CTGCCCCCTCAAGGGCGTGTGCGTACAGACACGACGCTGTTCTGACCGAAACTATCTG 120
DB 109 CTGCCCCCTCAAGGGCGTGTGCGTACAGACACGACGCTGTTCTGACCGAAACTATCTG 168
QY 121 AGGTCAACTTTCCTCCATTTTATTTAGGCAATTTCTTATGCTCCCATAGACGATGAG 180
DB 169 AGGTCAACTTTCCTCCATTTTATTTAGGCAATTTCTTATGCTCCCATAGACGATGAG 228
QY 181 AGCAGCAAGCAAGATGGGAATGACACCATTTGCTTCACTCTGTATAAAAAAGACG 240
DB 229 AGCAGCAAGCAAGATGGGAATGACACCATTTGCTTCACTCTGTATAAAAAAGACG 288
QY 241 GCCATGTGGGAGACCTTTCTGTGACGGGTGTGTGACAAAGAGATGATGCAAGAAATTAGA 300
DB 289 GCCATGTGGGAGACCTTTCTGTGACGGGTGTGTGACAAAGAGATGATGCAAGAAATTANA 348
QY 301 GAAAAATCTATTTTACAGACCAAGAGAGAGCAAGAAAGAGCTACAGAGCAAAAGCTGCA 360
DB 349 GAAAAATCTATTTTACAGACCAAGAGAGAGCAAGAAAGAGCTACAGAGCAAAAGCTGCA 408
QY 361 GCAAGCGGAGAGATCAAAATACCACTAGTGTGATGATGCAAGATTTGAA 411
DB 409 GCAAGCGGAGAGATCAAAATACCACTAGTGTGATGATGCAAGAGGAGAA 459

RESULT 15
US-09-918-995-10303
; Sequence 10303, Application US/09918995
; Publication No. US20030073623A1
; GENERAL INFORMATION:
; APPLICANT: Hyseq, Inc.
; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
; FILE REFERENCE: 20411-756
; CURRENT APPLICATION NUMBER: US/09/918,995
; CURRENT FILING DATE: 2001-07-30
; PRIOR APPLICATION NUMBER: US/09/235,076
; NUMBER OF SEQ ID NOS: 38054
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 10303
; LENGTH: 458
; TYPE: DNA
; ORGANISM: Homo sapiens
; NAME/KEY: misc_feature
; LOCATION: (1)...(458)
; OTHER INFORMATION: n = A,T,C or G
US-09-918-995-10303

Query Match      26.6%; Score 336; DB 10; Length 458;
Best Local Similarity 91.1%; Pred. No. 4e-60;
Matches 368; Conservative 0; Mismatches 35; Indels 1; Gaps 1;

QY 639 GAGAAATTCAGAAATATATTTACTGAGAAAGTTAAAGGAAGACAGTATTCCTGCTCG 698
DB 56 GAGAAAGACTGCAATATATTTACTGAGACGTTAAAGGAAGACAGTATTTACTGCACTCT 115
QY 699 CTCTGTTGGCAGTATTTAAATCACTTTACCCCTCGAGTATTTCCCAAGCTCTTCGTA 758
DB 116 GTGTGCTATGCGCTA-CCCCACCAACCCAGCTGAGAAATTTCCAATCAGTCTTCCTGA 174
QY 759 ATCACAAGTAGCAGAGAGGAGGAGTGCTACACAAACAAAGCTGAGGACCAAGAGCAAT 818
DB 175 ATTCAAGTAACACAGAGGAGGAGTGCTACACAAACAAAGCTGAGGACCAAGAGCAAT 234
QY 819 GAATCTGACATAGCTGAACCTTTGCGATTTTAAAGGAAGAAAGAAACCCAGCAATGGTT 878
DB 235 GAATCTGACATAGCTGAACCTTTGCGATTTTAAAGGAAGAAAGAAACCCAGCAATGGTT 294
QY 879 GAAGGATAAGGAAACAAATTTGTTTGCACCGGAAACCTATTTGGCAGCTATCAATGCATA 938
DB 295 GAAGGATAAGGAAACAAATTTGTTTGCACCGGAAACCTATTTGGCAGCTATCAATGCATA 354
QY 939 TAATTTAGCCATAGACTTAATTAATAGTCCCACTATTTGATTTGAACCGGCTGCTTG 998
DB 355 TAATTTAGCCATAGACTTAATTAATAGTCCCACTATTTGATTTGAACCGGCTGCTTG 414
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Qy 999 CCACCTAAACTAAAACTTACACAGGCTATTGAAGATTCTT 1042
Ob 415 CCACCTAAACTAAAACTTACACAGGCTATTGAAGATTCTT 458

Search completed: July 8, 2005, 07:35:57
Job time : 1946 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model
Run on: July 8, 2005, 09:23:13 ; Search time 861 Seconds
(without alignments)
9209.653 Million cell updates/sec

Title: US-10-681-199-1
Perfect score: 1263
Sequence: 1 atgctcttcaggtagcgca.....gaacagaactaaattcttaa 1263

Scoring table: OLIGO NUC
Gapop 60.0 , Gapext 60.0
Searched: 6330943 seqs, 3139157217 residues
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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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3	1263	100.0	1993	17	US-10-364-505-2
4	1263	100.0	1993	19	US-10-681-199-2
5	1002	79.3	1263	19	US-10-681-199-19
6	998	79.0	1559	17	US-10-108-260A-575
7	960	76.0	1263	19	US-10-681-199-13

Query Match 100.0%; Score 1263; DB 17; Length 1263;
Best Local Similarity 100.0%; Pred. No. 0;
Matches 1263; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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Db 1 ATGCTCTTCAGGTTAGCGATTACAGCTGGCAGCAGACGAGCTGGCTTTCTCTCT 60

ALIGNMENTS

RESULT 1
US-10-364-505-1
; Sequence 1, Application US/10364505
; Publication No. US20030219787A1
; GENERAL INFORMATION:
; APPLICANT: Kere, Juha
; APPLICANT: Taipale, Mikko
; APPLICANT: No. US20030219787A1ola-Hemmi, Jaana
; APPLICANT: Kaminen, Nina
; TITLE OF INVENTION: NOVEL HUMAN GENE FUNCTIONALLY RELATED TO DYSLEXIA
; FILE REFERENCE: 0933-0199P
; CURRENT APPLICATION NUMBER: US/10/364,505
; CURRENT FILING DATE: 2003-02-12
; NUMBER OF SEQ ID NOS: 13
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
; LENGTH: 1263
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: coding sequence for human DYXC1 (cDNA)
US-10-364-505-1

8 859 58.0 1263 19 US-10-681-199-15 Sequence 15, Appl
9 696 55.1 1263 19 US-10-681-199-17 Sequence 17, Appl
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c 11 325 25.7 325 18 US-10-240-425-872 Sequence 872, App
12 270 21.4 458 10 US-09-918-995-10303 Sequence 10303, A
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14 253 20.0 715 21 US-10-956-157-8349 Sequence 8349, Ap
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16 184 16.9 464 18 US-10-085-783A-30001 Sequence 30001, A
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RESULT 2
US-10-681-199-1
; Sequence 1, Application US/10681199
; Publication No. US20040138441A1
; GENERAL INFORMATION:
; APPLICANT: KERE, Juha
; TITLE OF INVENTION: NOVEL HUMAN GENE FUNCTIONALLY RELATED TO DYSLEXIA
; FILE REFERENCE: 0933-0214P
; CURRENT APPLICATION NUMBER: US/10/681,199
; CURRENT FILING DATE: 2003-10-09
; NUMBER OF SEQ ID NOS: 42
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 1
; LENGTH: 1263
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: coding sequence for human DYX1 (cDNA)
US-10-681-199-1

Query Match 100.0%; Score 1263; DB 19; Length 1263;
Best Local Similarity 100.0%; Pred. No. 0;
Matches 1263; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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; Sequence 2, Application US/10681199
; Publication No. US2004013841A1
; GENERAL INFORMATION:
; APPLICANT: KERE, Juha
; TITLE OF INVENTION: NOVEL HUMAN GENE FUNCTIONALLY RELATED TO DYSLEXIA
; FILE REFERENCE: 0933-0214P
; CURRENT APPLICATION NUMBER: US/10/681,199
; CURRENT FILING DATE: 2003-10-09
; NUMBER OF SEQ ID NOS: 42
; SOFTWARE: PatentIn ver. 2.1
; SEQ ID NO 2
; LENGTH: 1993
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (369)..(1628)
; FEATURE:
; OTHER INFORMATION: human DYXK1 mRNA as cDNA
US-10-681-199-2
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Best Local Similarity 100.0%; Pred. No. 0;
Matches 1263; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 909 TTATGTCAAAAAGAAAAAGCAAAATTAAGAAAGAGAGAAAAAATAAATAATTAAGAGTCTT 968
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Db 969 ACTAGAAATTTGGCATCTAGAAATCTTGCTCCAAAAGGGAGAAATTCAGAAAAATATATTT 1028
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Qy 721 AACTTTACCCCTCGAGTATTTCCCAACAGCTCTTGCTGTAATCACAACTAGCAGAGAGAG 780
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Qy 1261 TAA 1263
Db 1629 TAA 1631
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RESULT 5
US-10-681-199-19
; Sequence 19, Application US/10681199
; Publication No. US2004013841A1

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; GENERAL INFORMATION:
; APPLICANT: KERE, Juha
; TITLE OF INVENTION: NOVEL HUMAN GENE FUNCTIONALLY RELATED TO DYSLEXIA
; FILE REFERENCE: 0933-0214P
; CURRENT APPLICATION NUMBER: US/10/681.199
; CURRENT FILING DATE: 2003-10-09
; NUMBER OF SEQ ID NOS: 42
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 19
; LENGTH: 1263
; TYPE: DNA
; ORGANISM: Pan paniscus
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (1)..(1260)
; US-10-681-199-19

Query Match          79.3%; Score 1002; DB 19; Length 1263;
Best Local Similarity 99.6%; Pred. No. 0;
Matches 1252; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

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Db  7  CTTCAGGTTAGCGATTACAGCTGCGCAGACGAGCAAGACTGCGGTCTTCTGTCTCTCGCC 66

Qy  67  CTCAAAGCGGTGTCGTCAGAGACACGAGCTGTTCTGCACGGAATACTATCTGAAGTC 125
Db  67  CTCAAAGCGGTGTCGTCAGAGACACGAGCTGTTCTGCACGGAATACTATCTGAAGTC 126

Qy  127  AACTTTCTCCATTTTATTATTGAGGCATTTCTTTATGCTCCATAGACGATGAGAGCAGC 186
Db  127  AACTTTCTCCATTTTATTATTGAGGCATTTCTTTATGCTCCATAGACGATGAGAGCAGC 186

Qy  187  AAGCAAGAATTGGGAATGACACATTGTCTTCACTTTGATATAAAAGAGCGGCCATG 246
Db  187  AAGCAAGAATTGGGAATGACACATTGTCTTCACTTTGATATAAAAGAGCGGCCATG 246

Qy  247  TGGGAGACCTTTTCTGTGACGGGTGTTGACAAAGAGATGATGCAAGAATTAGAGAAAA 306
Db  247  TGGGAGACCTTTTCTGTGACGGGTGTTGACAAAGAGATGATGCAAGAATTAGAGAAAA 306

Qy  307  TCTATTTTACAAGCAAGAGAGACGCAAAAGAGCTACAGAAGCAAAAGCTGCAGCAAG 366
Db  307  TCTATTTTACAAGCAAGAGAGACGCAAAAGAGCTACAGAAGCAAAAGCTGCAGCAAG 366

Qy  367  CGGGAAGATCAAAATAACGACTAAGTGTGCATGATGAAGATTGAAGAAGAGAGAGAA 426
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Qy  427  AAAATGAGCATATGAAGAATAATGAACGGATAAAGCCACTAAAGCATTTGGAAGCTGG 486
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Qy  607  AATTGGCATCTAGAAATCTTTGCTCCAAAGGGAGAAATTCAGAAAAATATTTACTGAG 666
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Qy  667  AAGTTAAAGGAAGACAGTATTTCTGTCTCTCGCTCTGTGGCAGTATTAATAATCAACTTT 726
Db  667  AAGTTAAAGGAAGACAGTATTTCTGTCTCTCGCTCTGTGGCAGTATTAATAATCAACTTT 726

Qy  727  ACCCTCGAGTATTTCCCAACAGCTCTTTCGTGAATTCACAAGTAGCAGAGAGGAGTGG 786
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Db 283 GCCATGTGGGAGACCCCTTTCTGTGACGGGTGTTGACAAAGAGATGATGCAAGAAATTAGA 342
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Db 343 GAAAAATCTATTATTACAAGCACAAAGAGAGAGCAAAAAGAGCTACAGAAAGCAAAAGCTGCA 402
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Db 403 GCAAGCGGGAAGATCAAAATACGCACCTAAGTCTCATGTATGAAGATTGAAGAAGAGAG 462
Qy 421 AGGAAAAAATATAGAAATATCAAAAGAAATGAAAGGATGAAAGGCACTAAAGCAATTGGAA 480
Db 463 AGGAAAAAATATAGAAATATGAAAGAAATGAAAGGATGAAAGGCACTAAAGCAATTGGAA 522
Qy 481 GCCTGGAAGAAATATCAAAAGAAAGCTGAGAGCAAAAAAATTCAGAGAGAGAGAAA 540
Db 523 GCCTGGAAGAAATATCAAAAGAAAGCTGAGAGCAAAAAAATTCAGAGAGAGAGAAA 582
Qy 541 TTATGTCAAAAAGAAAGCAAAATTAAGAAGGAAGAAAGAAATATAAATATAAGAGTCTT 600
Db 583 TTATGTCAAAAAGAAAGCAAAATTAAGAAGGAAGAAAGAAATATAAATATAAGAGTCTT 642
Qy 601 ACTAGAAATTTGGCATCTAGAAATCTTGCTCCAAAGGGAGAAATTCAGAAAAATATATT 660
Db 643 ACTAGAAATTTGGCATCTAGAAATCTTGCTCCAAAGGGAGAAATTCAGAAAAATATATT 702
Qy 661 ACTGAGAGTTAAAGGAAGACAGTATTCCTGCTCCTCGCTCTGTTGGCAGTATTAAATC 720
Db 703 ACTGAGAGTTAAAGGAAGACAGTATTCCTGCTCCTCGCTCTGTTGGCAGTATTAAATC 762
Qy 721 AACTTTACCCCTCAGATTTCCCAACAGCTCTTCTGTAATCACAAGTAGCAGAGAGAG 780
Db 763 AACTTTACCCCTCAGATTTCCCAACAGCTCTTCTGTAATCACAAGTAGCAGAGAGAG 822
Qy 781 GAGTGGCTACACAAACAGCTGAGGCACGAAGCAATGAATCTGCATAGCTGAACTT 840
Db 823 GAGTGGCTACACAAACAGCTGAGGCACGAAGCAATGAATCTGCATAGCTGAACTT 882
Qy 841 TGCATTTAAAGGAAGAAAGAACCCGAAATGGTTGAAGGATAAAGGAACAAATG 900
Db 883 TGCATTTAAAGGAAGAAAGAACCCGAAATGGTTGAAGGATAAAGGAACAAATG 942
Qy 901 TTTGCAACGGAAACTATTGGCAGCTATCAATGCATATATTTAGCCATAAGACTAAAT 960
Db 943 TTTGCAACGGAAACTATTGGCAGCTATCAATGCATATATTTAGCCATAAGACTAAAT 1002
Qy 961 AATAAGATGCCATATTGTTGAAACCGGCTGCTTGGCCACCTTAAACTTAAACTTAA 1020
Db 1003 AATAAGATGCCATATTGTTGAAACCGGCTGCTTGGCCACCTTAAACTTAAACTTAA 1062
Qy 1021 CACAAGCTATTGAAGATTCTTCTAAGGC 1049
Db 1063 CACAAGCTATTGAAGATTCTTCTAAGGC 1091

RESULT 7
US-10-681-199-13
; Sequence 13, Application US/10681199
; Publication No. US20040138441A1
; GENERAL INFORMATION:
; APPLICANT: KERE, Juha
; TITLE OF INVENTION: NOVEL HUMAN GENE FUNCTIONALLY RELATED TO DYSLEXIA
; FILE REFERENCE: 0933-0214P
; CURRENT APPLICATION NUMBER: US/10/681.199
; CURRENT FILING DATE: 2003-10-09
; NUMBER OF SEQ ID NOS: 42
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 13
; LENGTH: 1263
; TYPE: DNA
; ORGANISM: Pan troglodytes
; FEATURE:
; NAME/KEY: CDS

; LOCATION: (1)..(1260)
US-10-681-199-13
Query Match 76.0%; Score 960; DB 19; Length 1263;
Best Local Similarity 99.6%; Pred. No. 0;
Matches 1210; Conservative 0; Mismatches 5; Indels 0; Gaps 0;
Qy 49 GTCTTTCTGTCTGTGCCCCCTCAAGGCGTGTGCGTCTAGACACGACGAGCTGTCTGTCACG 108
Db 49 GTCTTTCTGTCTGTGCCCCCTCAAGGCGTGTGCGTCTAGACACGACGAGCTGTCTGTCATG 108
Qy 109 GAAAACTATCTGAAGTCAACTTTCTCTCCATTTTATTTTGGGCAATTTCTTTATGCTCCC 168
Db 109 GAAAACTATCTGAAGTCAACTTTCTCTCCATTTTATTTTGGGCAATTTCTTTATGCTCCC 168
Qy 169 ATAGACGATGAGAGCAGCAAGCAAGATTGGGAATGACACCAATGTCTTCCCTTGAT 228
Db 169 ATAGACGATGAGAGCAGCAAGCAAGATTGGGAATGACACCAATGTCTTCCCTTGAT 228
Qy 229 AAAAAAGAGCGGCCCATGTGGGAGACCCCTTCTGTGACGGGTGTTGACAAAGAGATGATG 288
Db 229 AAAAAAGAGCGGCCCATGTGGGAGACCCCTTCTGTGACGGGTGTTGACAAAGAGATGATG 288
Qy 289 CAAAGAAATTAGAGAAAAATCTATTTTACAAGCACAGAGAGAGCAAAAGAGCTTACAGAA 348
Db 289 CAAAGAAATTAGAGAAAAATCTATTTTACAAGCACAGAGAGAGCAAAAGAGCTTACAGAA 348
Qy 349 GCAAAAGCTCAGCAAGCGGGAAGATCAAAAATACGCCATTAAGTGTCTATGATGAAGATT 408
Db 349 GCAAAAGCTCAGCAAGCGGGAAGATCAAAAATATGCACTAAGTGTCTATGATGAAGATT 408
Qy 409 GAAGAGAGAGAGGAAAAAATAGAGATATGAAGAAAAATGAACGGATATAAGCCATTAAGCCACT 468
Db 409 GAAGAGAGAGAGGAAAAAATAGAGATATGAAGAAAAATGAACGGATATAAGCCACT 468
Qy 469 AAAGCATTTGAAGCCTGGAAAGAAATATCAAGAAAAAGCTGAGGAGCAAAAAAATTCAG 528
Db 469 AAAGCATTTGAAGCCTGGAAAGAAATATCAAGAAAAAGCTGAGGAGCAAAAAAATTCAG 528
Qy 529 AGAGAGAGAAATATGTCAAAAAGAAAGCAAAATTAAGAGAGAGCAAAAAAATTAATA 588
Db 529 AGAGAGAGAAATATGTCAAAAAGAAAGCAAAATTAAGAGAGAGCAAAAAAATTAATA 588
Qy 589 TATAAGAGTCTTACTAGAAATTTGGCATCTAGAAATCTTCTCCAAAGGAGAGAAATCA 648
Db 589 TATAAGAGTCTTACTAGAAATTTGGCATCTAGAAATCTTCTCCAAAGGAGAGAAATCA 648
Qy 649 GAAAAATATTTACTGAGAAAGTTAAAGGAAGACAGTATTCCTGCTCCTGCTGTGTTGGC 708
Db 649 GAAAAATATTTACTGAGAAAGTTAAAGGAAGACAGTATTCCTGCTCCTGCTGTGTTGGC 708
Qy 709 AGTATTAATAATCAACTTTTACCCCTCGAGTATTCCAAACAGCTCTTCTGTAATCAAGTA 768
Db 709 AGTATTAATAATCAACTTTTACCCCTCGAGTATTCCAAACAGCTCTTCTGTAATCAAGTA 768
Qy 769 GCAGAGAGGAGGAGTGGCTACACAAACAGCTGAGGAGAGCAAGCAAGCAAGTAACTGAC 828
Db 769 GCAGAGAGGAGGAGTGGCTACACAAACAGCTGAGGAGAGCAAGCAAGTAACTGAC 828
Qy 829 ATAGCTGAACTTTTCCGATTTTAAAGGAAGAAAGAAACCCAGAAATGTTGAAGGATATA 888
Db 829 ATAGCTGAACTTTTCCGATTTTAAAGGAAGAAAGAAACCCAGAAATGTTGAAGGATATA 888
Qy 889 GGAACAAATTTGTTGCAACGGAATAATTTTGGCAGCTATCAATGATATTAATTTAGCC 948
Db 889 GGAACAAATTTGTTGCAACGGAATAATTTTGGCAGCTATCAATGATATTAATTTAGCC 948
Qy 949 ATAAGACTTAATAATAGATGCCACTATTTGATTTGAAACCGGGCTGCTTGGCCACTTAAA 1008
Db 949 ATAAGACTTAATAATAGATGCCACTATTTGATTTGAAACCGGGCTGCTTGGCCACTTAAA 1008
Qy 1009 CTAAAAAACTTTACAAAGGCTATTGAAGATTCTTCTAAGGCAGCTGGAATTTATGATGCA 1068
Db 1009 CTAAAAAACTTTACAAAGGCTATTGAAGATTCTTCTAAGGCAGCTGGAATTTATGATGCA 1068

Db	1009	CTAAAAA	CTTACACAAGG	CTATTGAAGATTTCTTCTAAGGCAC	CTGGAATTTATTGATGCCA	1068
Qy	1069	CCTGTTACAGACA	CAATGCTTAATGCAAGAA	TGAAGGCACATGTACGACGTGGAACAGCA	TTCC	1128
Db	1069	CCTGTTACAGACA	CAATGCTTAATGCAAGAA	TGAAGGCACATGTACGACGTGGAACAGCA	TTCC	1128
Qy	1129	TGTCAACTAGAA	TTGTTATGTAGAAGG	CCTACAGGATTTAGAGCGGCAC	TTTAAGATTGAT	1188
Db	1129	TGTCAACTAGAA	TTGTTATGTAGAAGG	CCTACAGGATTTAGAGCGGCAC	TTTAAGATTGAT	1188
Qy	1189	CCATCCAA	CAAAAATTTGACAAA	ATGATGCTGAGAAAGATTCGGAA	TTGAAATTCAGGAA	1248
Db	1189	CCATCCAA	CAAAAATTTGACAAA	ATGATGCTGAGAAAGATTCGGAA	TTGAAATTCAGGAA	1248
Qy	1249	GAACTAAA	ATCTTAA	1263		
Db	1249	GAACTAAA	ATCTTAA	1263		

RESULT 8

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US-10-681-199-15
; Sequence 15, Application US/10681199
; Publication No. US20040138441A1
; GENERAL INFORMATION:
; APPLICANT: KERE, Juha
; TITLE OF INVENTION: NOVEL HUMAN GENE FUNCTIONALLY RELATED TO DYSLEXIA
; FILE REFERENCE: 0933-0214P
; CURRENT APPLICATION NUMBER: US/10/681,199
; CURRENT FILING DATE: 2003-10-09
; NUMBER OF SEQ ID NOS: 42
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 15
; LENGTH: 1263
; TYPE: DNA
; ORGANISM: Gorilla gorilla
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (1)..(1260)
US-10-681-199-15

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[illegible]

RESULT 9

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; TITLE OF INVENTION: NOVEL HUMAN GENE FUNCTIONALLY RELATED TO DYSLEXIA
; SEQUENCE ID: 10681199
; APPLICATION NO.: US20040138441A1
; PUBLICATION NO.: US20040138441A1
; GENERAL INFORMATION:
; APPLICANT: KERE, Juhua
; FILING DATE: 2003-10-09
; CURRENT FILING DATE: 2003-10-09
; CURRENT APPLICATION NUMBER: US/10/681,199
; NUMBER OF SEQ ID NOS: 42
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 17
; LENGTH: 1263
; TYPE: DNA
; ORGANISM: Pongo pygmaeus
; FEATURE:

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; NAME/KEY: CDS
; LOCATION: (1)...(1260)
US-10-681-199-17

Query Match      55.1%; Score 696; DB 19; Length 1263;
Best Local Similarity 99.1%; Pred. No. 0;
Matches 1246; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

Qy 7 CTTTCAGTTAGCGATTACAGTGGCAGCAGACGAGAGTGGTCTTCTGCTCTGCCC 66
Db 7 CTTTCAGTTAGCGATTACAGTGGCAGCAGACGAGAGTGGTCTTCTGCTCTGCCC 66

Qy 67 CTCAAGCGGTGTCGTCAGAGACGAGAGTGGTCTGACGAGAACTATCTGAAGTC 126
Db 67 CTCAAGCGGTGTCGTCAGAGACGAGAGTGGTCTGACGAGAACTATCTGAAGTC 126

Qy 127 AACTTTCTCATTTTATTTAGGCAATTTCTTATGCTCCCATAGACGATGAGAGCAGC 186
Db 127 AACTTTCTCATTTTATTTAGGCAATTTCTTATGCTCCCATAGACGATGAGAGCAGC 186

Qy 187 AAAGCAAGATTGGGAATGACACCAATTTCTTCACTTGTATATAAAGAGCGCCATG 246
Db 187 AAAGCAAGATTGGGAATGACACCAATTTCTTCACTTGTATATAAAGAGCGCCATG 246

Qy 247 TGGGAGACCCCTTTCTGTGACGGGTGTTGACAAAGAGACGATGCAAGAAATTTAGAGAAA 306
Db 247 TGGGAGACCCCTTTCTGTGACGGGTGTTGACAAAGAGACGATGCAAGAAATTTAGAGAAA 306

Qy 307 TCTATTTTACAAGCACAAGAGAGAGCAAAAGAGCTACAGAGCAAAAGCTGCAAGCAAG 366
Db 307 TCTATTTTACAAGCACAAGAGAGAGCAAAAGAGCTACAGAGCAAAAGCTGCAAGCAAG 366

Qy 367 CGGGAAGATCAAAATACGCAATTAAGTGTCTGATGATGAAGATTGAAGAAAGAGAGAAA 426
Db 367 CGGGAAGATCAAAATATGCAATTAAGTGTCTGATGATGAAGATTGAAGAAAGAGAGAAA 426

Qy 427 AAAATAGAGATGAAAGAAATGAAAGGATGAAAGGATGAAAGGATGAAAGGATGAAAG 486
Db 427 AAAATAGAGATGAAAGAAATGAAAGGATGAAAGGATGAAAGGATGAAAGGATGAAAG 486

Qy 487 AAAGAAATATCAAGAAAGCTGAGAGCAAAAGAAATTTAGAGAGAGAGAAATTTATGT 546
Db 487 AAAGAAATATCAAGAAAGCTGAGAGCAAAAGAAATTTAGAGAGAGAGAAATTTATGT 546

Qy 547 CAAAAAGAAAGCAAAATTAAGAAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 606
Db 547 CAAAAAGAAAGCAAAATTAAGAAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 606

Qy 607 AATTGGCATCTAGAAATCTTGTCCAAAGAGGAGAAATTCAGAGAAATATTTACTGAG 666
Db 607 AATTGGCATCTAGAAATCTTGTCCAAAGAGGAGAAATTCAGAGAAATATTTACTGAG 666

Qy 667 AAGTTAAAGAGAGACAGTATTTCTGCTCTGCTCTGCTCTGCTCTGCTCTGCTCTGCT 726
Db 667 AAGTTAAAGAGAGACAGTATTTCTGCTCTGCTCTGCTCTGCTCTGCTCTGCTCTGCT 726

Qy 727 ACCCTCGAGTATTTCCCAAGCTCTTGTGTAATCAAGATGAGAGAGAGAGAGAGAG 786
Db 727 ACCCTCGAGTATTTCCCAAGCTCTTGTGTAATCAAGATGAGAGAGAGAGAGAGAG 786

Qy 787 CTACACAAAGCTGAGGACGAGAGCAATGATGATGATGATGATGATGATGATGATGAT 846
Db 787 CTACACAAAGCTGAGGACGAGAGCAATGATGATGATGATGATGATGATGATGATGAT 846

Qy 847 TTAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 906
Db 847 TTAAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 906

Qy 907 ACAGAAACTATTTGGAGCTATCAATGATATATTTAGCCATTAAGCTAAATAAAG 966
Db 907 ACAGAAACTATTTGGAGCTATCAATGATATATTTAGCCATTAAGCTAAATAAAG 966

Qy 967 ATGCCACTATTTGATTTGAACGGGCTGCTTGGCCACCTTAAACTTAAAGAACTTACACAAG 1026
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Db 967 ATGCCACTATTTGATTTGAACGGGCTGCTTGGCCACCTTAAAGAACTTACACAAG 1026
Qy 1027 GCTATTGAAGATTTCTTAAAGGACCTGGAATTAATGATGACCACTGTTACAGACAATGCT 1086
Db 1027 GCTATTGAAGATTTCTTAAAGGACCTGGAATTAATGATGACCACTGTTACAGACAATGCT 1086
Qy 1087 AATGCAAGATTAAGGACACATGATGACGCTGGAAACAGCATTTCTGTAACCTAGAAATGCTAT 1146
Db 1087 AATGCAAGATTAAGGACACATGATGACGCTGGAAACAGCATTTCTGTAACCTAGAAATGCTAT 1146
Qy 1147 GTAGAAGGCTTACAGGATTAAGGCGGCACTTAAAGATTGATCCATCCAAACAAATTTGTA 1206
Db 1147 GTAGAAGGCTTACAGGATTAAGGCGGCACTTAAAGATTGATCCATCCAAACAAATTTGTA 1206
Qy 1207 CAAATTTGATGCTGAGAAAGATTCGGAATGTAATTTCAAGGAAACAGAACTTAAATCTTAA 1263
Db 1207 CAAATTTGATGCTGAGAAAGATTCGGAATGTAATTTCAAGGAAACAGAACTTAAATCTTAA 1263

RESULT 10
US-09-918-995-22675
; Sequence 22675, Application US/09918995
; Publication No. US20030073623A1
; GENERAL INFORMATION:
; APPLICANT: Hyseq, Inc.
; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
; FILE REFERENCE: 20411-756
; CURRENT APPLICATION NUMBER: US/09/918,995
; PRIOR FILING DATE: 2001-07-30
; PRIOR APPLICATION NUMBER: US/09/235,076
; NUMBER OF SEQ ID NOS: 38054
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 22675
; LENGTH: 488
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-918-995-22675

Query Match      25.8%; Score 326; DB 10; Length 488;
Best Local Similarity 100.0%; Pred. No. 1.7e-151;
Matches 326; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 80 GCGTCAGAGACACGACGCTGTTCTGACGGAACCTATCTGAAGGTCACTTCTCCAT 139
Db 73 GCGTCAGAGACACGACGCTGTTCTGACGGAACCTATCTGAAGGTCACTTCTCCAT 132

Qy 140 TTTTATTTGAGGCAATTTCTTATGCTCCCATAGACGATGAGAGCAAGAAAGATTG 199
Db 133 TTTTATTTGAGGCAATTTCTTATGCTCCCATAGACGATGAGAGCAAGAAAGATTG 192

Qy 200 GGAATGACACCAATTTCTTCACTTGTATATAAAGAGCGGCCATGTTGGAGACCTTTT 259
Db 193 GGAATGACACCAATTTCTTCACTTGTATATAAAGAGCGGCCATGTTGGAGACCTTTT 252

Qy 260 CTGTGACGGGTGTTGACAAAGAGATGATGCAAGAAATAGAGAAATCTATTTTACAAG 319
Db 253 CTGTGACGGGTGTTGACAAAGAGATGATGCAAGAAATAGAGAAATCTATTTTACAAG 312

Qy 320 CACAAGAGAGAGCAAAAGAGCTACAGAGCAAAAGCTGCGAGCAAGCGGAGAGATCAA 379
Db 313 CACAAGAGAGAGCAAAAGAGCTACAGAGCAAAAGCTGCGAGCAAGCGGAGAGATCAA 372

Qy 380 AATACGCACTAAGTGTCTATGATGAAG 405
Db 373 AATACGCACTAAGTGTCTATGATGAAG 398

RESULT 11
US-10-425-872/c
; Sequence 872, Application US/10240425
```

Publication No. US20040033502A1

GENERAL INFORMATION:
APPLICANT: Williams, Amanda
APPLICANT: Bolland, Joseph F.
APPLICANT: Lord, Reginald V.
APPLICANT: Alvarez, Chris
APPLICANT: Wetzel, Jon C.
APPLICANT: Scherff, Uwe
APPLICANT: Vockley, Joseph G.
TITLE OF INVENTION: Gene Expression Profiles in Esophageal Tissue
FILE REFERENCE: 44921-5026
CURRENT APPLICATION NUMBER: US/10/240,425
CURRENT FILING DATE: 2002-09-30
PRIOR APPLICATION NUMBER: PCT/US01/09847
PRIOR FILING DATE: 2001-03-28
PRIOR APPLICATION NUMBER: US 60/193,446
PRIOR FILING DATE: 2000-03-31
NUMBER OF SEQ ID NOS: 1588
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 872
LENGTH: 325
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
OTHER INFORMATION: Genbank Accession No. US20040033502A1 AI783611
US-10-240-425-872

Query Match 25.7%; Score 325; DB 18; Length 325;
Best Local Similarity 100.0%; Pred. No. 5.2e-151;
Matches 325; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 524 TTCAGAGAGAGAGAAATATGTCAAAAGAAAAGCAAAATTAAGAGAGAGAGAAAAA 583
DB 325 TTCAGAGAGAGAGAAATATGTCAAAAGAAAAGCAAAATTAAGAGAGAGAGAAAAA 266

QY 584 TAAATATAAGAGCTTACTAGAAATTTGGCAATCTAGAAATCTGCTCCAAAGGAGAA 643
DB 265 TAAATATAAGAGCTTACTAGAAATTTGGCAATCTAGAAATCTGCTCCAAAGGAGAA 206

QY 644 ATTCAAGAAATATATTACTGAGAGAGTTAAAGGAGAGAGATTTCTGCTCTCTCTG 703
DB 205 ATTCAAGAAATATATTACTGAGAGTTAAAGGAGAGAGATTTCTGCTCTCTCTG 146

QY 704 TTGGCAGTATTAATCAACTTTACCTCGAGTATTCCTCAACAGCTTTCTGTGAATCAC 763
DB 145 TTGGCAGTATTAATCAACTTTACCTCGAGTATTCCTCAACAGCTTTCTGTGAATCAC 86

QY 764 AAGTAGCAGAGAGAGAGGAGTGCTTACACAAACAGCTGAGCAGAGAGCGAATGAATA 823
DB 85 AAGTAGCAGAGAGAGAGGAGTGCTTACACAAACAGCTGAGCAGAGAGCGAATGAATA 26

QY 824 CTGACATAGCTGAACCTTTGCGATTT 848
DB 25 CTGACATAGCTGAACCTTTGCGATTT 1

RESULT 12

US-09-918-995-10303
Sequence 10303, Application US/09918995
Publication No. US20030073623A1
GENERAL INFORMATION:
APPLICANT: Hyseq, Inc.
TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
FROM VARIOUS CDNA LIBRARIES
FILE REFERENCE: 20411-756
CURRENT APPLICATION NUMBER: US/09/918,995
CURRENT FILING DATE: 2001-07-30
PRIOR APPLICATION NUMBER: US/09/235,076
PRIOR FILING DATE: 1999-01-20
NUMBER OF SEQ ID NOS: 38054
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 10303
LENGTH: 458

TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: misc feature
LOCATION: (1)...(458)
OTHER INFORMATION: n = A,T,C or G
US-09-918-995-10303

Query Match 21.4%; Score 270; DB 10; Length 458;
Best Local Similarity 100.0%; Pred. No. 1.4e-123;
Matches 270; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 773 AAGAGGAGAGTGGCTACACAAACAGCTTGAGGACGAGAGCAATGATCTGACATAG 832
DB 189 AAGAGGAGAGTGGCTACACAAACAGCTTGAGGACGAGAGCAATGATCTGACATAG 248

QY 833 CTGAACCTTTGGCATTTAAAGAGAGAGAAAGCAACCCAGAGATGGTTGAAGGATAAGGAA 892
DB 249 CTGAACCTTTGGCATTTAAAGAGAGAGAGAAAGCAACCCAGAGATGGTTGAAGGATAAGGAA 308

QY 893 ACAAATTTGTCACACGAGAACTATTTGGCAGCTATCAATGCATATATTTAGGCATAA 952
DB 309 ACAAATTTGTCACACGAGAACTATTTGGCAGCTATCAATGCATATATTTAGGCATAA 368

QY 953 GACTAAATATAAGATGCCACTATTTGTAATTTGAACCGGCTGCTTGCACCTTAAACTAA 1012
DB 369 GACTAAATATAAGATGCCACTATTTGTAATTTGAACCGGCTGCTTGCACCTTAAACTAA 428

QY 1013 AAACTTACACAGGCTATTGAAGATCTT 1042
DB 429 AAACTTACACAGGCTATTGAAGATCTT 458

RESULT 13

US-10-956-157-3114/c
Sequence 3114, Application US/10956157
Publication No. US20050118625A1
GENERAL INFORMATION:
APPLICANT: Wyeth
APPLICANT: Mounts, William
TITLE OF INVENTION: NUCLEIC ACID ARRAYS FOR DETECTING GENE EXPRESSION ASSOCIATED WITH
HUMAN OSTEOARTHRITIS AND HUMAN PROTEASES
FILE REFERENCE: 031896-043000 (AM 101081)
CURRENT APPLICATION NUMBER: US/10/956,157
CURRENT FILING DATE: 2004-10-04
NUMBER OF SEQ ID NOS: 319805
SOFTWARE: PatentIn version 3.2
SEQ ID NO 3114
LENGTH: 715
TYPE: DNA
ORGANISM: Homo sapiens
US-10-956-157-3114

Query Match 20.0%; Score 253; DB 21; Length 715;
Best Local Similarity 100.0%; Pred. No. 4.2e-115;
Matches 253; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 638 GGAGAAATTCAGAAAAATATTTACTGAGAGAGTTAAAGGAGAGACAGTATTTCTGCTCTC 697
DB 430 GGAGAAATTCAGAAAAATATTTACTGAGAGTTAAAGGAGAGACAGTATTTCTGCTCTC 371

QY 698 GCTCTGTTGGCAGTATTAATAATCAACTTTACCCCTCGAGTATTCCTCAACAGCTTTCTG 757
DB 370 GCTCTGTTGGCAGTATTAATAATCAACTTTACCCCTCGAGTATTCCTCAACAGCTTTCTG 311

QY 758 AATCAAGTAGCAGAGAGAGGAGTGCTTACACAAACAGCTGAGCAGAGAGAGAA 817
DB 310 AATCAAGTAGCAGAGAGAGGAGTGCTTACACAAACAGCTGAGCAGAGAGAGAA 251

QY 818 TGAATCTGACATAGCTGAACCTTTGCGATTTTAAAGAGAGAGAGAAAGAACCCAGATG 877
DB 250 TGAATCTGACATAGCTGAACCTTTGCGATTTTAAAGAGAGAGAGAAAGAACCCAGATG 191

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OM nucleic - nucleic search, using sw model

Run on: July 8, 2005, 06:20:46 ; Search time 9619 Seconds
(without alignments)
4997.939 Million cell updates/sec

Title: US-10-681-199-1

Perfect score: 1263

Sequence: 1 atgctcttcaggtagcgca.....gaacagaactaaatcttaa 1263

Scoring table: OLIGO NUC

Gapop 60.0 , Gapext 60.0

Searched: 34239544 seqs, 19032134700 residues

Word size : 0

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database :

EST:*
1: gb_est1:*
2: gb_est2:*
3: gb_hic:*
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9: gb_gse2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	838	66.3	1600	3	BC017392	BC017392 Homo sapi
2	583	46.2	844	6	CD107587	CD107587 AGENCOURT
3	462	36.6	535	7	CN478982	CN478982 UI-CF-FNO
4	446	35.3	1131	5	B0217312	B0217312 AGENCOURT
5	422	33.4	462	7	CN429788	CN429788 170006000
6	410	32.5	788	2	BE564350	BE564350 601343161
7	405	32.1	823	5	B0567908	B0567908 AGENCOURT
8	378	29.9	792	4	BG771796	BG771796 602720472
9	374	29.6	933	6	CD358543	CD358543 AGENCOURT
10	364	28.8	573	2	BE972748	BE972748 601652170
11	342	27.1	772	6	CA422987	CA422987 UI-H-FLO-
12	340	26.9	468	6	CB297042	CB297042 12B22058
13	338	26.8	417	1	A1024221	A1024221 ov81e02.s
14	325	25.7	325	1	A1783611	A1783611 tz99c01.x
15	323	25.6	651	5	B0852580	B0852580 AGENCOURT
16	320	25.3	701	4	BG482634	BG482634 602502783
17	313	24.8	1183	2	BF207765	BF207765 601861861
18	290	23.0	665	4	BG540324	BG540324 602568825
19	275	21.8	559	5	BU607404	BU607404 UI-CF-FNO
20	274	21.7	625	2	BF216970	BF216970 601884034
21	251	19.9	645	5	BM984145	BM984145 UI-CF-DU1
22	250	19.8	270	2	BF372375	BF372375 PM4-PT002
23	244	19.3	744	4	BG192162	BG192162 RST11269
24	241	19.1	267	1	A1073572	A1073572 ov45g11.x

c 25	218	17.3	583	6	CB160627	CB160627 K-EST0220
c 26	216	17.1	382	1	AA724419	AA724419 ah91f07.s
c 27	215	17.0	775	4	BG206612	BG206612 RST26063
c 28	212	16.8	523	2	BE178005	BE178005 RC3-HT060
c 29	197	15.6	251	1	AA317003	AA317003 EST18868
c 30	182	14.4	458	2	BE463906	BE463906 hy18f08.x
c 31	175	13.9	988	4	BI517373	BI517373 603041624
c 32	150	11.9	752	4	BG183140	BG183140 RST2158.A
c 33	148	11.7	940	4	BG220651	BG220651 RST40439
c 34	146	11.6	855	2	BF248143	BF248143 601859338
c 35	105	8.3	467	1	AI674107	AI674107 wd18c04.x
c 36	102	8.1	500	4	BI517981	BI517981 603041624
c 37	85	6.7	459	5	EX095509	EX095509 BX095509
c 38	78	6.2	402	2	BE178258	BE178258 RC3-HT060
c 39	78	6.2	683	5	BP460416	BP460416 BP460416
c 40	70	5.5	112	4	BM820152	BM820152 K-EST0088
c 41	64	5.1	440	1	AI360851	AI360851 qx99h12.x
c 42	61	4.8	793	7	CO738958	CO738958 SILR04c21
c 43	60	4.8	654	5	BM971229	BM971229 UI-CF-DU1
c 44	57	4.5	515	5	BU738973	BU738973 UI-E-EJ0-
c 45	51	4.0	285	8	AQ067717	AQ067717 HS_2239_B

ALIGNMENTS

RESULT 1
BC017392
LOCUS Homo sapiens, Similar to RIKEN cDNA 1700010I24 gene, clone
DEFINITION IMAGE:4081622, mRNA.
ACCESSION BC017392
VERSION BC017392.1 GI:19263480
KEYWORDS HTC.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 1600)
AUTHORS Strausberg,R.
TITLE Direct Submission
JOURNAL Submitted (13-NOV-2001) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-4590, USA
REMARK NIH-MGC Project URL: <http://mgc.nci.nih.gov>
COMMENT Contact: MGC help desk
Email: cgapbs-remail.nih.gov
Tissue Procurement: ATCC
cDNA Library Preparation: CLONTECH Laboratories, Inc.
DNA Sequencing by: The I.M.A.G.E. Consortium (LLNL)
<http://www.sysgenbiology.org>
contact: amadaneysystemsbiology.org
Anup Madan, Jessica Fahey, Erin Helton, Mark Kettelman, Anuradha Madan, Stephanie Rodriguez, Amy Sanchez and Michelle Whiting

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>
Series: IRAL Plate: 32 Row: k Column: 22
This clone was selected for full length sequencing because it passed the following selection criteria: Hexamer frequency ORF analysis, GenomeScan gene prediction
This clone has the following problem: frame shifted.

FEATURES
source
Location/Qualifiers
1..1600
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:4081622"
/tissue_type="Bladder, carcinoma"
/clone_lib="NIH_MGC_53"
/lab_host="DH10B"

cDNA Library Preparation: Life Technologies, Inc.
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
 DNA Sequencing by: Agencourt Bioscience Corporation
 Clone distribution: MGC clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
 Plate: LLAMI3293 row: n column: 13
 High quality sequence stop: 388.
 Location/Qualifiers
 1. .1131
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:6047052"
 /tissue_type="melanotic melanoma"
 /lab_host="DH10B (phage-resistant)"
 /clone_lib="NIH_MGC_72"
 /notes="Organ: skin; Vector: pCMV-SPORT6; Site 1: NotI;
 Site 2: SalI; Cloned unidirectionally. Primer: Oligo dt.
 Average insert size 2 kb. Library constructed by Life
 Technologies."

FEATURES

source

ORIGIN
 Query Match 35.3%; Score 446; DB 5; Length 1131;
 Best Local Similarity 100.0%; Pred. No. 9.1e-217; Indels 0; Gaps 0;
 Matches 446; Conservative 0; Mismatches 0;

Qy 42 GACTGCGGTCTTTCTGTCCTGCCCTCAAGGCGTGTGCAGACACGACGCGTGT 101

Db 1 GACTGCGGTCTTTCTGTCCTGCCCTCAAGGCGTGTGCAGACACGACGCGTGT 60

Qy 102 CTGCACGGAACATCTGAGGTCACCTTCTCCATTTTATTGAGGCATTTCTTTA 161

Db 61 CTGCACGGAACATCTGAGGTCACCTTCTCCATTTTATTGAGGCATTTCTTTA 120

Qy 162 TGCTCCCATAGACATGAGACGACCAAGCAAGATGGGAATGACACCATTTCTTCAC 221

Db 121 TGCTCCCATAGACATGAGACGACCAAGCAAGATGGGAATGACACCATTTCTTCAC 180

Qy 222 CTTGTATAAAAGAACGCGCATGTGGAGACCCCTTCTGTGACGGGTGTGACAAGA 281

Db 181 CTTGTATAAAAGAACGCGCATGTGGAGACCCCTTCTGTGACGGGTGTGACAAGA 240

Qy 282 GATGATCGAAGATTAGAGAAATCTATTTTACAGCACAGAGAGCAAGAGC 341

Db 241 GATGATCGAAGATTAGAGAAATCTATTTTACAGCACAGAGAGCAAGAGC 300

Qy 342 TACAGAGCAAAAGCTGCAGCAAGCGGAGATCAAAAATACGCATTAAGTGTATGAT 401

Db 301 TACAGAGCAAAAGCTGCAGCAAGCGGAGATCAAAAATACGCATTAAGTGTATGAT 360

Qy 402 GAAGATTGAAGAACAGAGAGGAAAAAATAGAGATATGAAGAAATGAACGGATAAA 461

Db 361 GAAGATTGAAGAACAGAGAGGAAAAAATAGAGATATGAAGAAATGAACGGATAAA 420

Qy 462 AGCCACTAAGCATTGGAGCCTGGA 487

Db 421 AGCCACTAAGCATTGGAGCCTGGA 446

RESULT 5
 CN429788
 LOCUS CN429788 462 bp mRNA linear EST 16-MAY-2004
 DEFINITION 1700600026401 GRN_PREHEP Homo sapiens cDNA 5', mRNA sequence.
 ACCESSION CN429788
 VERSION CN429788.1 GI:47417382

KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 462)
 AUTHORS Brandenberger, R., Wei, H., Zhang, S., Lei, S., Murage, J., Fisk, G.J.,

TITLE

JOURNAL

COMMENT

Li, Y., Xu, C., Fang, R., Guegler, K., Rao, M.S., Mandalam, R.,
 Lebkowski, J. and Stanton, L.W.
 Transcriptome characterization elucidates signaling networks that
 control human ES cell growth and differentiation
 Nat. Biotechnol. 22 (6), 707-716 (2004)
 Contact: Brandenberger R
 Regenerative Medicine
 Geron Corporation
 230 Constitution Drive, Menlo Park, CA 94025, USA
 Tel: 650 473 8658
 Fax: 650 473 7760
 Email: rbrandenberger@geron.com
 Insert Length: 462 Std Error: 0.00.

FEATURES

source

1. .462
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /tissue_type="embryonic stem cells, DMSO-treated H9 cell
 line"
 /clone_lib="GRN_PREHEP"
 /notes="Oligo dt primed, full-length enriched cDNA library
 from DMSO-treated HES cell line H9 (p22) maintained in
 feeder-free conditions"

ORIGIN

Query Match 33.4%; Score 422; DB 7; Length 462;
 Best Local Similarity 100.0%; Pred. No. 1.7e-204; Indels 0; Gaps 0;
 Matches 422; Conservative 0; Mismatches 0;

Qy 1 ATGCTCTTTCAGGTAGCGATTACAGCTGGCAGCAGACGAAAGACTGCGGTCTTTCTGTCT 60

Db 41 ATGCTCTTTCAGGTAGCGATTACAGCTGGCAGCAGACGAAAGACTGCGGTCTTTCTGTCT 100

Qy 61 CTGCCCTCAAAAGCGGTGCGTCAGAGACACGACGCTGTTGCGAGGAAAATCTATCTG 120

Db 101 CTGCCCTCAAAAGCGGTGCGTCAGAGACACGACGCTGTTGCGAGGAAAATCTATCTG 160

Qy 121 AAGGTCAACTTCTCCATTTTATTGAGGCATTTCTTATGTCCTCCATAGACGATGAG 180

Db 161 AAGGTCAACTTCTCCATTTTATTGAGGCATTTCTTATGTCCTCCATAGACGATGAG 220

Qy 181 AGCAGCAAAAGCAAGATTGGGAATGACACCATTTGTCTTCACTTGTATATAAAGAGCG 240

Db 221 AGCAGCAAAAGCAAGATTGGGAATGACACCATTTGTCTTCACTTGTATATAAAGAGCG 280

Qy 241 GCATGTGGAGACCCCTTCTGTGACGGGTGTGACAAAGAGATGATGCAAGAAATTAGA 300

Db 281 GCATGTGGAGACCCCTTCTGTGACGGGTGTGACAAAGAGATGATGCAAGAAATTAGA 340

Qy 301 GAAAAATCTATTTTACAAGCACAAAGAGAGCAAAAGAACTACAGAAAGCAAGCTGCA 360

Db 341 GAAAAATCTATTTTACAAGCACAAAGAGAGCAAAAGAACTACAGAAAGCAAGCTGCA 400

Qy 361 GCAAAGCGGGAAGATCAAAAATACGCATTAAGTGTCTATGATGAAGATTGAAGAGAAAGAG 420

Db 401 GCAAAGCGGGAAGATCAAAAATACGCATTAAGTGTCTATGATGAAGATTGAAGAGAAAGAG 460

Qy 421 AG 422

Db 461 AG 462

RESULT 6
 BE564350
 LOCUS BE564350 788 bp mRNA linear EST 15-AUG-2000
 DEFINITION 601343161F1 NIH_MGC_53 Homo sapiens cDNA clone IMAGE:3685335 5',
 mRNA sequence.
 ACCESSION BE564350
 VERSION BE564350.1 GI:9808070
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens


```

QY 361 GCAAAGCGGAAGATCAAAATATACGACTAAGTGTCAATGATGAAG 405
Db 412 GCAAAGCGGAAGATCAAAATATACGACTAAGTGTCAATGATGAAG 456

RESULT 8
BG771796
LOCUS 792 bp mRNA linear EST 15-MAY-2001
DEFINITION 602720472F1 NIH_MGC_97 Homo sapiens cDNA clone IMAGE:4837505 5',
mRNA sequence.
ACCESSION BG771796
VERSION BG771796.1 GI:14082449
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 792)
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
TITLE Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
JOURNAL NIH-MGC http://mgi.nci.nih.gov/.
COMMENT National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
CDNA Library Preparation: Michael J. Brownstein (NHGRI), Shiraki
Toshiyuki and Piero Carninci (RIKEN)
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Incyte Genomics, Inc.
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: LHAM10769 row: p column: 18
High quality sequence stop: 671.
FEATURES
source
Location/Qualifiers
1..792
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:4837505"
/lab_host="DH10B"
/clone_lib="NIH_MGC 97"
/notes="Organ: testis; Vector: pBluescriptR (modified
pBluescript KS+); Site 1: BamHI; Site 2: SalI-XhoI
(gtcgag); Oligo-dT primed using primer
5'-TTTTTTTTTTTTTTVN-3', size-selected for average
insert size 2.2 kb and normalized to ROT 5. This is a
primary library enriched for full-length clones and
constructed using the Cap-trapper method (Carninci, in
preparation)." Library constructed by M. Brownstein
(NIH/NHGRI, National Institutes of Health). Note: this is
a NIH_MGC Library."

ORIGIN
Query Match 29.9%; Score 378; DB 4; Length 792;
Best Local Similarity 99.8%; Pred. No. 6 4e-182;
Matches 428; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ATGCCTCTTCAGGTACGATTACAGCTGGCAGCAGACGAACTGCGGTCTTCTGCT 60
Db 37 ATGCCTCTTCAGGTACGATTACAGCTGGCAGCAGACGAACTGCGGTCTTCTGCT 96

QY 61 CTGCCCTCAAGCGGTGCGTCAGACACGACGCTGTCACGGAACATATCTG 120
Db 97 CTGCCCTCAAGCGGTGCGTCAGACACGACGCTGTCACGGAACATATCTG 156

QY 121 AAGGTCAACTTCTCCATTTTATTTAGGCACTTCTTATGCTCCCATAGCATGAG 180
Db 157 AAGGTCAACTTCTCCATTTTATTTAGGCACTTCTTATGCTCCCATAGCATGAG 216

QY 181 AGCAGCAAGCAAGATTGGGAATGACACCATTTCTTCACTTGTATATAAAGAAAGCG 240
Db 217 AGCAGCAAGCAAGATTGGGAATGACACCATTTCTTCACTTGTATATAAAGAAAGCG 276

QY 241 GCCATGTGGAGACCCCTTTCTGTGACGGGTGTTGACAAAGAGAGATGATGCAAAAGATTAGA 300
Db 277 GCCATGTGGAGACCCCTTTCTGTGACGGGTGTTGACAAAGAGATGATGCAAAAGATTAGA 336

QY 301 GAAAAATCTATTTTACAAGCACAAGAGAGAGCAAAAGAGAGCTTACAGAACGAAAAGCTGCA 360
Db 337 GAAAAATCTATTTTACAAGCACAAGAGAGAGCAAAAGAGAGCTTACAGAACGAAAAGCTGCA 396

QY 361 GCAAAGCGGGAGATCAAAATATACGACTAAGTGTCTATGATGAAGAGAGAGAG 420
Db 397 GCAAAGCGGGAGATCAAAATATACGACTAAGTGTCTATGATGAAGAGAGAGAGAG 456

QY 421 AGCAAAAAA 429
Db 457 AGCAAAAAA 465

RESULT 9
CD358543
LOCUS 933 bp mRNA linear EST 29-MAY-2003
DEFINITION AGENCOURT_14255831 NIH_MGC_180 Homo sapiens cDNA clone
IMAGE:30386203 5', mRNA sequence.
ACCESSION CD358543
VERSION CD358543.1 GI:31129978
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 933)
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
TITLE Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
JOURNAL NIH-MGC http://mgi.nci.nih.gov/.
COMMENT National Institutes of Health, Mammalian Gene Collection (MGC)
Unpublished (1999)
Contact: Daniela S. Gerhard, Ph.D.
Office of Cancer Genomics
National Cancer Institute / NIH
Bldg. 31 Rm10A07 Bethesda, MD 20892
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: Dr. Michael Brownstein
CDNA Library Preparation: Invitrogen Corp
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
Plate: NDAM454 row: h column: 20
High quality sequence start: 303
High quality sequence stop: 666.
FEATURES
source
Location/Qualifiers
1..933
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:30386203"
/lab_host="DH10B-Ton A ( T1 and T5 phage resistances)"
/clone_lib="NIH_MGC 180"
/notes="Organ: Testis; Vector: pCMV-SPORT6.1; Site 1: NotI;
Site 2: EcoRV (destroyed); Library is oligo-dT primed and
directionally cloned (EcoRV site is destroyed upon
cloning). Average insert size 1.68 kb. Library was
constructed by (Invitrogen). Note: this is a NIH_MGC
Library."

ORIGIN
Query Match 29.6%; Score 374; DB 6; Length 933;
Best Local Similarity 99.8%; Pred. No. 7.3e-180;
Matches 424; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 437 ATATGAAGAAATGAACGATAAAGCCACTAAGCATTTGGAGCCTGGAAAGATATC 496
Db 313 ATATGAAGAAATGAACGATAAAGCCACTAAGCATTTGGAGCCTGGAAAGATATC 372

```



```
/lab host="DH10B (Life Technologies)"
/clone_lib="NCI CGAP_FLO"
/notes="Organ: Chondrosarcoma; Vector: pT7T3-Pac
(Pharmacia) with a modified polylinker; Site 1: EcoR I;
Site 2: Not I; NCI CGAP FLO is a cDNA library derived from
a pool of mRNA obtained from 4 cell lines from grade III
chondrosarcoma tissues. The library was constructed
according to Bonaldo, Lennon and Soares, Genome Research,
6:791-806, 1996. First strand cDNA synthesis was primed
with an oligo-dT primer containing a Not I site. Double
stranded cDNA was ligated to an EcoR I adaptor, digested
with Not I, and cloned directionally into pT7T3-Pac
vector. The oligonucleotide used to prime the synthesis of
first-strand cDNA contains a library tag sequence that is
located between the Not I site and the (dT)18 tail. The
sequence tag for this library is GAGTCGGTG. The cell line
was provided by Dr James Martin from University of Iowa.
TAG TISSUE=Human Chondrosarcoma Grade 3 cell line mix
TAG_LIB=UI-H-FLO
TAG_SEQ=GAGTCGGTG"
```

ORIGIN

```
Query Match      27.1%; Score 342; DB 6; Length 772;
Best Local Similarity 100.0%; Pred. No. 1.8e-163; Indels 0; Gaps 0;
Matches 342; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 922 GCAGCTATCATATATAATTTAGCCATAAGACTAAATAATAAGATGCCACTATTGTAT 981
Db 720 GCAGCTATCATATATAATTTAGCCATAAGACTAAATAATAAGATGCCACTATTGTAT 661

Qy 982 TTGAACCGGGCTGCTGCCACCTAAACTAAAACTTACCAAGGCTATTGAAGATTCT 1041
Db 660 TTGAACCGGGCTGCTGCCACCTAAACTAAAACTTACCAAGGCTATTGAAGATTCT 601

Qy 1042 TCTAAGCACTGGAATTATTGATGCCACCTGTTACAGCAATGCTAATGCAAGAAATGAAG 1101
Db 600 TCTAAGCACTGGAATTATTGATGCCACCTGTTACAGCAATGCTAATGCAAGAAATGAAG 541

Qy 1102 GCACATGTACGACGTGGAACAGCACTTCTGCACTAGAAATGTATGTAAGAGGCTACAG 1161
Db 540 GCACATGTACGACGTGGAACAGCACTTCTGCACTAGAAATGTATGTAAGAGGCTACAG 481

Qy 1162 GATTATGAGCGGCACCTTAAGATTGATCCATCCACAAATTTGACAAATTTGATGCTGAG 1221
Db 480 GATTATGAGCGGCACCTTAAGATTGATCCATCCACAAATTTGACAAATTTGATGCTGAG 421

Qy 1222 AAGATTTCGGATGTAATTTCAAGGAACAGAACTAAATCTTAA 1263
Db 420 AAGATTTCGGATGTAATTTCAAGGAACAGAACTAAATCTTAA 379
```

```
RESULT 12
CB297042      468 bp      mRNA      linear      EST 28-FEB-2003
LOCUS      12B22058_rev_1_F09_r_075.ab1 Chimpanzee brain library Koo's Pan
DEFINITION      troglodytes cDNA clone 12B22058_rev_1_F09_r_075.ab1 5', mRNA
sequence.
CB297042
CB297042.1      GI:28622472
VERSION      EST.
KEYWORDS      Pan troglodytes (chimpanzee)
SOURCE      Pan troglodytes
ORGANISM      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
REFERENCE      1 (bases 1 to 468)
AUTHORS      Hellmann,I., Zollner,S., Enard,W., Ebersberger,I., Nickl,B. and
Paabo,S.
TITLE      Selection on human genes as revealed by comparisons to chimpanzee
cDNA
JOURNAL      Genome Res. (2003) In press
COMMENT      Contact: Paabo S
Evolutionary Genetics
Max-Planck-Institute for evolutionary Anthropology
```

```
Deutscher Platz 6, 04103 Leipzig, Germany
Tel: +49 (0)-341-3550 500
Fax: +49 (0)-341-3550 555
Email: paabo@eva.mpg.de
Seq primer: M13 reverse.
FEATURES
Location/Qualifiers
source      1..468
            /organism="Pan troglodytes"
            /mol_type="mRNA"
            /db_xref="taxon:9598"
            /clone="12B22058_rev_1_F09_r_075.ab1"
            /sex="male"
            /tissue type="brain, presumably cortex"
            /dev stage="adult"
            /lab_host="Epichurian Coli (TM) XL-10-Gold"
            /clone_lib="Chimpanzee brain library Koo's"
            /notes="Vector: pUCHi; Site 1: SfiI-A; Site 2: SfiI-B; The
            library was prepared using the SMART cDNA library
            construction kit (Clontech), doing only primer extension,
            but not PCR amplification of the cDNA. The only deviation
            from the published protocol was that we cloned the cDNA
            into a plasmid vector."
```

```
ORIGIN
Query Match      26.9%; Score 340; DB 6; Length 468;
Best Local Similarity 99.7%; Pred. No. 1.9e-162; Indels 0; Gaps 0;
Matches 390; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 49 GTCTTTCTGTCTCTGCCCTCAAGGCGTGTGGTCTAGAGACACGACGCGTGTCTGCACG 108
Db 78 GTCTTTCTGTCTCTGCCCTCAAGGCGTGTGGTCTAGAGACACGACGCGTGTCTGCACG 137

Qy 109 GAAACTATCTGAAGTCAACTTTCTCCATTTTATTTAGGAGCATTTCTTATGCTCCC 168
Db 138 GAAACTATCTGAAGTCAACTTTCTCCATTTTATTTAGGAGCATTTCTTATGCTCCC 197

Qy 169 ATAGACGATGAGAGCAGCAAGCAAGATTTGGGAATGACACCATTTGTCCTCACCTTGAT 228
Db 198 ATAGACGATGAGAGCAGCAAGCAAGATTTGGGAATGACACCATTTGTCCTCACCTTGAT 257

Qy 229 AAAAAAGAGCGGCATGTGGGAGACCCCTTTCTGTGCGGTGTGTGACAAAGAGATGATG 288
Db 258 AAAAAAGAGCGGCATGTGGGAGACCCCTTTCTGTGCGGTGTGTGACAAAGAGATGATG 317

Qy 289 CAAAGATTAGAGAAAATCTATTTTACAGACACAGAGAGAGCAAAAGAGCTACAGAA 348
Db 318 CAAAGATTAGAGAAAATCTATTTTACAGACACAGAGAGAGCAAAAGAGCTACAGAA 377

Qy 349 GCAAAAGCTGCAGCAAGCGGGAAGATCAAAAATACCACTAAGTGTGATGATGAAGATT 408
Db 378 GCAAAAGCTGCAGCAAGCGGGAAGATCAAAAATACCACTAAGTGTGATGATGAAGATT 437

Qy 409 GAAGAAGAGAGAGAGAAAAAATAGAAAGATA 439
Db 438 GAAGAAGAGAGAGAGAAAAAATAGAAAGATA 468
```

```
RESULT 13
AI024221      417 bp      mRNA      linear      EST 18-JUN-1998
LOCUS      ov81e02.sl Soares_testis_NHT Homo sapiens cDNA clone IMAGE:1643738
DEFINITION      3', mRNA sequence.
AI024221
AI024221.1      GI:3239265
VERSION      EST.
KEYWORDS      Homo sapiens (human)
SOURCE      Homo sapiens
ORGANISM      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE      1 (bases 1 to 417)
AUTHORS      NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE      National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
```


JOURNAL
COMMENT

Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-re@mail.nih.gov
cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima Bonaldo, Ph.D.
cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Seq primer: -40m13 fwd. ET from Amersham.

FEATURES
source
1..417
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:1643738"
/sex="male"
/lab_host="DH10B"
/clone_lib="Soares testis_NHT"
/note="Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA was prepared from mRNA obtained from Clontech Laboratories, Inc., and primed with a Not I - oligo (dT) primer [5',
TGTTACCAATCTGAAGTGGAGCGCGCCCAATTTTTTTTTTTTTT 3'].
Double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Not I and cloned into the Not I and Eco RI sites of the modified pT73 vector. Library went through one round of normalization to Cot5, and was constructed by Bento Soares and M. Fatima Bonaldo."

ORIGIN
Query Match 26.8%; Score 338; DB 1; Length 417;
Best Local Similarity 99.7%; Pred. No. 2e-161;
Matches 388; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 149 AGGCATTTCTTTATGCTCCATAGACGATGAGAGCAGCAAGCAAGATTGGGAATGACA 208
Db 1 AGGCATTTCTTTATGCTCCATAGACGATGAGAGCAGCAAGCAAGATTGGGAATGACA 60
Qy 209 CCATGTCTTCACCTGTATATAAAGAGCGGCCATGTGGAGACCTTTCTGTGACGG 268
Db 61 CCATGTCTTCACCTGTATATAAAGAGCGGCCATGTGGAGACCTTTCTGTGACGG 120
Qy 269 GTGTTGACAAAGAGATGATGCAAGCAATTAGAGAAAATCTATTTTACAGCACAAGAGA 328
Db 121 GTGTTGACAAAGAGATGATGCAAGCAATTAGAGAAAATCTATTTTACAGCACAAGAGA 180
Qy 329 GAGCAAAAAGAGCTTACAGAGCAAAAGCTGCAAGCAAGCGGGAAGATCAAAAATACGCAC 388
Db 181 GAGCAAAAAGAGCTTACAGAGCAAAAGCTGCAAGCAAGCGGGAAGATCAAAAATACGCAC 240
Qy 389 TAAGTGTCTATGATGAAGATTGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 448
Db 241 TAAGTGTCTATGATGAAGATTGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAGAG 300
Qy 449 ATGAACGATATAAGCCACTAAAGCATTTGGAGCCTGGAAGAAATATCAAGAGAAAGCTG 508
Db 301 ATGAACGATATAAGCCACTAAAGCATTTGGAGCCTGGAAGAAATATCAAGAGAAAGCTG 360
Qy 509 AGGAGCAAAAAGAGATTTTACAGAGAGAGAGAG 537
Db 361 AGGAGCAAAAAGAGATTTTACAGAGAGAGAGAG 389

RESULT 14
AI783611/c
LOCUS
DEFINITION
similar to contains 'Alu repetitive element', mRNA sequence.
ACCESSION
AI783611.1
VERSION
AI783611.1
GI:5325420

KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

EST.
Homo sapiens (human)
Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 325)
NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
Unpublished (1997)
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-re@mail.nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R. Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Insert Length: 437 Std Error: 0.00
Seq primer: -40UP from Gibco
High quality sequence stop: 325.
Location/Qualifiers
1..325
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:2296704"
/lab_host="DH10B"
/clone_lib="NCI-CGAP_Kid11"
/note="Organ: kidney; Vector: pT73D-Pac (Pharmacia) with a modified polylinker; Site 1: Not I; Site 2: Eco RI; Plasmid DNA from the normalized library NCI CGAP Kid3 was prepared, and ss circles were made in vitro. Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (cloneIDs 1322376-1323911, 1456007-1456775, and 1500552-1502855). Subtraction by Bento Soares and M. Fatima Bonaldo."

ORIGIN
Query Match 25.7%; Score 325; DB 1; Length 325;
Best Local Similarity 100.0%; Pred. No. 8.9e-155;
Matches 325; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
Qy 524 TTCAGAGAGAGAGAAATTTATGTCAAAAGAAAAGCAAAATTAAGAGAGAGAGAAAAA 583
Db 325 TTCAGAGAGAGAGAAATTTATGTCAAAAGAAAAGCAAAATTAAGAGAGAGAGAAAAA 266
Qy 584 TAAATATAAGAGCTTTACTAGAAATTTGGCATCTAGAAATCTTGTCTCCTCGCTCTG 643
Db 265 TAAATATAAGAGCTTTACTAGAAATTTGGCATCTAGAAATCTTGTCTCCTCGCTCTG 206
Qy 644 ATTGAGAAAAATATATTTTACTGAGAGAGTTAAAGAGAGAGAGATTTCTGCTCTGCTG 703
Db 205 ATTGAGAAAAATATATTTTACTGAGAGAGTTAAAGAGAGAGAGATTTCTGCTCTGCTG 146
Qy 704 TTGCGAGTATTAATCAACTTTACCTTCCCTCGAGTATTTCCCAACAGCTTTTGTGATCAC 763
Db 145 TTGCGAGTATTAATCAACTTTTACCTTCCCTCGAGTATTTCCCAACAGCTTTTGTGATCAC 86
Qy 764 AAGTAGCAG 823
Db 85 AAGTAGCAG 26
Qy 824 CTGACATAGCTGAACCTTTTGGGATTT 848
Db 25 CTGACATAGCTGAACCTTTTGGGATTT 1

RESULT 15

BUS52580
LOCUS BUS52580 651 bp mRNA linear EST 16-OCT-2002
DEFINITION AGENCOURT_10501009 NIH_MGC_82 Homo sapiens cDNA clone IMAGE:6619012
5', mRNA sequence.
ACCESSION BUS52580
VERSION BUS52580.1 GI:24037543
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 651)
AUTHORS NIH-MGC <http://mgc.nci.nih.gov/>.
TITLE National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL Unpublished (1999)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgapbs-r@mail.nih.gov
Tissue Procurement: CLONTECH
CDNA Library Preparation: CLONTECH Laboratories, Inc.
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
<http://image.llnl.gov>
Plate: LCM2866 row: n column: 04
High quality sequence stop: 364.
FEATURES
source
1..651
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:6619012"
/lab_host="DH10B (T1 phage-resistant)"
/clone_lib="NIH_MGC_82"
/note="Organ: Testis; Vector: pDNR-LIB (Clontech); Site: 1:
SfiI (ggccattatggcc); Site 2: SfiI (ggccattatggcc); 5' and
3' adaptors were used in cloning as follows: 5' adaptor
sequence: 5'-CAGCGCATTATGCGC-3' and 3' adaptor sequence:
5'-ATTCTAGCGCGAGCGCGGCGACATG-dt(30)BN-3' (where B = A,
C, or G and N = A, C, G, or T). Average insert size
1.35 kb (range 0.9-4.0 kb). 14/15 colonies contained
inserts by PCR. This library was enriched for full-length
clones and was constructed by Clontech Laboratories (Palo
Alto, CA)."
ORIGIN
Query Match 25.6%; Score 323; DB 5; Length 651;
Best Local Similarity 99.7%; Pred. No. 1e-153;
Matches 373; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 875 GGTTGAAGGATAAAGGAACAATAATTGTTGCAACGGAAACTATTGGCAGCTATCAATG 934
Db 3 GGTTGAAGGATAAAGGAACAATAATTGTTGCAACGGAAACTATTGGCAGCTATCAATG 62
Qy 935 CATATAATTTAGCCATAGACTAATAATAAGATGCCACTATTGTATTGAACCGGCTG 994
Db 63 CATATAATTTAGCCATAGACTAATAATAAGATGCCACTATTGTATTGAACCGGCTG 122
Qy 995 CTTCGCCACCTAAAACTAAAAAATTACACAAGGCTATTGAAGATTCCTTAAGGCACCTGG 1054
Db 123 CTGCACCTAAAACTAAAAAATTACACAAGGCTATTGAAGATTCCTTAAGGCACCTGG 182
Qy 1055 AATTATTGATGCCACCTGTTACAGACATGCTAATGCAAGATGAAGGCACATGTACGAC 1114
Db 183 AATTATTGATGCCACCTGTTACAGACATGCTAATGCAAGATGAAGGCACATGTACGAC 242
Qy 1115 GTGGAACAGCAATTCGTCAACTAGAAATTTGTATGTAGAGGCTACAGGATTTATCAAGCGG 1174
Db 243 GTGGAACAGCAATTCGTCAACTAGAAATTTGTATGTAGAGGCTACAGGATTTATCAAGCGG 302
Qy 1175 CACTTAAGATTGATCCATCCAAACAAATTTGACAAATTTGATGCTGAGAAGATTCGGAATG 1234
Db 303 CACTTAAGATTGATCCATCCAAACAAATTTGACAAATTTGATGCTGAGAAGATTCGGAATG 362

Qy 1235 TAATTCAAGGAACA 1248
|||||
Db 363 TAATTCAAGGAACA 376
Search completed: July 8, 2005, 12:03:37
Job time : 9626 secs